

DESCRIPTION

THERMOPLASTIC ELASTOMER COMPOSITION
AND MOLDED ARTICLE

5

TECHNICAL FIELD

The present invention relates to a thermoplastic elastomer composition excellent in moldability, heat resistance, weather resistance, chemical resistance, adhesivity, flexibility and abrasion resistance. Further, the present invention relates to a composition for powder slash molding and a powder slash molded article using the composition.

BACKGROUND ART

15 It is known that an acrylic block copolymer having methyl methacrylate as a hard segment and butyl acrylate as a soft segment has properties as a thermoplastic elastomer. For example, the mechanical properties of an acrylic block copolymer having a methacryl block and an acryl block which is produced by an iniferter method are disclosed (Patent Reference 1).

20 The acrylic block copolymer has characteristics excellent in weather resistance, heat resistance, durability, oil resistance and abrasion resistance. Further, an elastomer which is extremely soft in comparison with other thermoplastic elastomers such as a styrene block copolymer can be provided by suitably selecting components composing the block copolymer.

As uses making use of the properties of the acrylic block

copolymer, there has been expected development as various superficial skin materials, interior materials, and a material for a part which is directly touched by a hand making use of smooth feeling thereof.

As physical properties necessary for these superficial skin materials, there are resistance to chemicals which may be possibly contacted, further, the adhesivity of superficial skin with a substrate when the superficial skin and the substrate are directly adhered, the adhesivity of the superficial skin with a buffer material when the buffer material is provided between the superficial skin and the substrate, in addition to mechanical properties, scratch resistance, heat resistance, strain restorability. As the molding method of the superficial skin materials, the powder slash molding using soft powder material which is a powder molding method is widely adopted for the superficial skin of interior equipments for an automobile such as an instrument panel, a console box and a door trim. This is caused by that they have soft texture, skin crepe and stitch can be provided, further, the degree of freedom in design is large, and designing property is satisfactory. Since a forming pressure is not applied in the molding method, differing from other molding methods such as injection molding and compression molding, it is a condition that powder flowability is excellent because the powder material is required to be uniformly adhered on a mold with a complicated shape at molding and simultaneously, it is also a condition that melt viscosity is low because powder adhering on a mold is melted and a film is required to be formed by flowing under no pressure. As the material, a polyvinyl chloride sheet has been conventionally widely used because it is excellent in hardness of a surface in use and flexibility, but since a

polyvinyl chloride resin contains a lot of chlorine in its molecules, there is fear of great adverse effect on environments and an effective substitute material is desired (Patent Reference 2). Accordingly, a sheet molded article of a thermoplastic elastomer has been
5 conventionally developed as the substitution of the polyvinyl chloride resin (Patent Reference 3, Patent Reference 4, and Patent Reference 5). However, sheets using a polyolefin resin and a styrene elastomer are insufficient in abrasion resistance, flexibility and oil resistance. Further, a sheet using thermoplastic polyurethane was poor in
10 moldability, and also has a problem from the viewpoint of the cost thereof.

Patent Reference 1: Japanese Patent No. 2553134

Patent Reference 2: JP-A-5-279485

Patent Reference 3: JP-A-7-82433

15 Patent Reference 4: JP-A-10-30036

Patent Reference 5: JP-A-2000-103957

DISCLOSURE OF INVENTION

An object of the present invention is to obtain an acrylic
20 block copolymer composition improving melt flowability at molding and being excellent in heat resistance in addition to maintain weather resistance, chemical resistance, adhesivity, flexibility and abrasion resistance which are the characteristics of the acrylic block copolymer.

As a result of repeating intensive studies in order to solve
25 the above-mentioned problems, the present inventors have found that the problems can be effectively solved by converting the acrylic block copolymer to having a high molecular weight or crosslinking the acrylic

block copolymer at molding, and reached the present invention.

Namely, the present invention relates to a thermoplastic elastomer composition comprising an acrylic block copolymer (A) which comprises a methacrylic polymer block (a) and an acrylic polymer block (b), wherein at least one of polymer blocks among the methacrylic polymer block (a) and the acrylic polymer block (b) has a functional group (X), and a compound (B) containing 1.1 or more of functional groups (Y) in one molecule.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein the functional group (X) is at least one kind of functional groups selected from an acid anhydride group, a carboxyl group, a hydroxyl group and an epoxy group.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein the functional group (Y) is at least one kind of functional groups selected from an epoxy group, a carboxyl group, a hydroxyl group, an amino group, an acid anhydride group and an oxazoline group.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein the functional group (X) is an acid anhydride group and/or a carboxyl group, and the functional group (Y) is an epoxy group.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein a boiling point of the compound (B) is 200°C or more.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein the compound (B) is a polymer having the weight average molecular weight of 50,000 or less.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein the acrylic block copolymer (A) comprises 10 to 60 % by weight of the methacrylic polymer block (a) in which a methacrylic polymer is the main component and 90 to 40 % by weight of the acrylic polymer block (b) in which the acrylic polymer is the main component.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein the acrylic polymer block (b) comprises 50 to 100 % by weight of at least one monomer selected from the group consisting of n-butyl acrylate, ethyl acrylate and 2-methoxyethyl acrylate and 50 to 0 % by weight of other acrylate and/or other vinyl monomer copolymerizable with these monomers.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein the number average molecular weight of the acrylic block copolymer (A) measured by gel permeation chromatography is 30,000 to 200,000.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein a ratio (M_w/M_n) of the weight average molecular weight to the number average molecular weight measured by gel permeation chromatography of the acrylic block copolymer (A) is 1.8 or less.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein the acrylic block copolymer (A) is a block copolymer produced by atom transfer radical polymerization.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein a glass transition temperature of the methacrylic polymer block (a) is 25 to 130°C.

A preferable embodiment relates to the thermoplastic elastomer composition, wherein 5 to 200 parts by weight of a filler is further added based on 100 parts by weight of the acrylic block copolymer (A).

5 A preferable embodiment relates to the thermoplastic elastomer composition, wherein 0.1 to 20 parts by weight of a lubricant is further added based on 100 parts by weight of the acrylic block copolymer (A).

Also, the present invention relates to the thermoplastic
10 elastomer composition for powder slash molding, containing the above-mentioned composition.

The present invention relates to a molded article, which is obtained by powder slash molding the above-mentioned composition.

The present invention relates to a superficial skin for an
15 automobile interior, which is obtained by powder slash molding the above-mentioned composition.

EFFECT OF THE INVENTION

An acrylic block copolymer composition excellent in
20 moldability and heat resistance in addition to maintain weather resistance, chemical resistance, adhesivity, flexibility and abrasion resistance, which are the characteristics of an acrylic block copolymer, can be obtained by using the present invention. Further, the composition of the present invention can be preferably used for powder
25 slash molding.

BEST MODE FOR CARRYING OUT THE INVENTION

The present invention is further explained in detail in the following.

The present invention is characterized in comprising a thermoplastic elastomer composition comprising an acrylic block copolymer (A) which comprises a methacrylic polymer block (a) and an acrylic polymer block (b), wherein at least one of polymer blocks among the methacrylic polymer block (a) and the acrylic polymer block (b) has a functional group (X), and a compound (B) containing at least 1.1 or more of functional groups (Y) in one molecule. The functional group (Y) is a functional group having reactivity with the functional group (X) and usually reacted at molding, whereby the acrylic block copolymer (A) is converted to having a high molecular weight or crosslinked.

<Acrylic block copolymer (A)>

The structure of the acrylic block copolymer (A) of the present invention is a linear block copolymer or a branched (star shape) block copolymer and a mixture thereof. The structure of the block copolymer may be used depending on the physical properties of the acrylic block copolymer (A) required, but the linear block copolymer is preferable from the viewpoints of cost and easiness of polymerization.

The linear block copolymer may be any structure. When the methacrylic polymer block (a) is represented as "a" and the acrylic polymer block (b) is represented as "b", it is preferable that at least one acrylic block copolymer selected from the group consisting of (a-b)_n type, b-(a-b)_n type and (a-b)_n-a type (n is an integer of at least 1, for example, an integer of 1 to 3), from the viewpoint of the physical

properties of the linear block copolymer or the physical properties of the composition. It is not particularly limited, but among those, an a-b type diblock copolymer, an a-b-a type triblock copolymer, or a mixture thereof is preferable from the viewpoints of easiness of handling at processing and the physical properties of the composition.

It is characterized in that at least one of the functional group (X) is introduced in at least one polymer block of the methacrylic polymer block (a) and the acrylic polymer block (b). When its number is at least 2, the mode in which the monomer is polymerized may be random copolymerization or block copolymerization. When the a-b-a type triblock copolymer is represented as an example, any of a (a/z)-b-a type, a (a/z)-b-(a/z) type, a z-a-b-a type, a z-a-b-a-z type, an a-(b/z)-a type, an a-b-z-a type, an a-z-b-z-a type, a (a/z)-(b/z)-(a/z) type, and a z-a-z-b-z-a-z type may be satisfactory. Herein, z represents a monomer or a polymer block containing the reactive functional group (X), (a/z) represents that a monomer containing the functional group (X) is copolymerized in the methacrylic polymer block (a) and (b/z) represents that a monomer containing the functional group (X) is copolymerized in the acrylic polymer block (b).

Further, a site containing z and a mode of containing z in the methacrylic polymer block (a) or the acrylic polymer block (b) may be freely set, and can be used in accordance with a purpose.

The number average molecular weight of the acrylic block copolymer (A) is not particularly limited, and may be determined from molecular weight respectively necessary for the methacrylic polymer block (a) and the acrylic polymer block (b). When the molecular weight is small, there may be a case where mechanical properties

enough as an elastomer cannot be expressed, and adversely when the molecular weight is large more than necessity, there may be a case where processing property is lowered. Since flowage under non pressurization is required in particular in case of the powder slash
5 molding, when the molecular weight is large, melt viscosity is enhanced and moldability tends to be deteriorated.

From the above-mentioned viewpoint, the number average molecular weight measured by gel permeation chromatography of the acrylic block copolymer (A) is preferably 30,000 to 200,000, more
10 preferably 35,000 to 150,000, and further preferably 40,000 to 100,000.

The ratio (M_w/M_n) of the weight average molecular weight to the number average molecular weight measured by gel permeation chromatography of the acrylic block copolymer (A) is not particularly
15 limited, but is preferably 1.8 or less and more preferably 1.5 or less. When M_w/M_n exceeds 1.8, there may be a case where uniformity of the acrylic block copolymer is deteriorated.

The composition ratio of the methacrylic polymer block (a) to the acrylic polymer block (b) which form the acrylic block copolymer
20 (A) is 5 to 90 % by weight of the block (a) and 95 to 10 % by weight of the block (b). The range of the composition ratio is preferably 10 to 60 % by weight of (a) and 90 to 40 % by weight of (b), and further preferably 15 to 50 % by weight of (a) and 85 to 50 % by weight of (b) from the viewpoints of retention of a shape at molding and elasticity as
25 an elastomer. When the proportion of (a) is less than 5 % by weight, the shape tends to be hardly retained at molding and when the proportion of (b) less than 10 % by weight, elasticity as an elastomer

and melting property at molding tend to be lowered.

Since when the proportion of (a) is small, hardness tends to be low and when the proportion of (b) is small, hardness tends to be high, from the viewpoint of hardness of the elastomer composition, the proportion can be set according to the hardness of the required elastomer composition. Further, since when the proportion of (a) is small, viscosity tends to be low, and when the proportion of (b) is small, viscosity tends to be high, from the viewpoint of the processing, the proportion can be set according to the required processing property.

The relation of glass transition temperatures between the methacrylic polymer block (a) and the acrylic polymer block (b) which form the acrylic block copolymer (A) preferably satisfies the relation of the following formula, referring to the glass transition temperature of the methacrylic polymer block (a) as T_{ga} and referring to the glass transition temperature of the acrylic polymer block (b) as T_{gb} .

$$T_{ga} > T_{gb}$$

The setting of the glass transition temperature (T_g) of the above-mentioned polymer (the methacrylic polymer block (a) and acrylic polymer block (b)) can be carried out by setting the weight ratio of a monomer at respective polymer portions according to the following Fox formula.

$$\begin{aligned} 1/T_g &= (W_1/T_{g1}) + (W_2/T_{g2}) + \dots + (W_m/T_{gm}) \\ W_1 + W_2 + \dots + W_m &= 1 \end{aligned}$$

Wherein T_g represents the glass transition temperature of a polymer portion and T_{g1} , T_{g2} , ---, T_{gm} represent the glass transition temperatures of respective polymerization monomers. Further, W_1 , W_2 , ----, W_m represent weight ratios of respective polymerization
5 monomers.

As the glass transition temperatures of respective polymerization monomers in the above-mentioned Fox formula, for example, values described in Polymer Handbook Third Edition (Wiley - Interscience, 1989) may be used.

10 Further, the glass transition temperature can be measured by DSC (Differential Scanning Calorimeter) or the $\tan \delta$ peak of dynamic viscoelasticity, but when polarity between the methacrylic polymer block (a) and the acrylic polymer block (b) is too near and the chain number of block monomers is too little, there may be a case
15 where those measurement values are deviated from the calculation formula by the above-mentioned Fox formula.

<Reactive functional group (X)>

In the present invention, the functional group (X) is not particularly limited, it is enough to act as a reaction point with the
20 compound (B), preferably acts as a reaction point for converting the acrylic copolymer (A) to having a high molecular weight or a crosslinking point for crosslinking the acrylic block copolymer (A), and is preferably at least one kind of functional groups selected from an acid anhydride group, a carboxyl group, a hydroxyl group and an
25 epoxy group from the viewpoints of the stability of a bond prepared by the reaction, the easiness of the reaction, the easiness of introduction to the acrylic block copolymer (A), and cost thereof. Among them, at

least one kind of functional groups selected from an acid anhydride group and a carboxyl group is more preferable from the viewpoints of the easiness of introduction to the acrylic block copolymer (A) and the cost thereof.

5 These functional groups (X) are introduced into the acrylic block copolymer (A) by a form protecting the functional group with an appropriate protecting group or the form of the precursors of the functional group (X) and then, the functional group (X) can be also prepared by known chemical reactions.

10 These functional groups (X) can be used in a combination of two or more kinds thereof, and when two or more kinds are used in combination, it is preferable to select functional groups which are not mutually reacted.

 The functional group (X) may be contained in either of the
15 methacrylic polymer block (a) or the acrylic polymer block (b), may be contained in both blocks, and can be used so that the introduction condition of the functional group (X) is preferable in accordance with purposes such as the reaction point of the acrylic block copolymer (A), the cohesive force and glass transition temperature of the block (the
20 methacrylic polymer block (a) and the acrylic polymer block (b)) composing the acrylic block copolymer (A) and further the required physical property of the acrylic block copolymer (A).

 For example, when the methacrylic polymer block (a) and the acrylic polymer block (b) are wanted to be selectively reacted with
25 the functional group (X) as reaction points using the compound (B) containing the functional group (Y) which has reactivity with the functional group (X), the functional group (X) may be introduced in

blocks to be reacted.

Further, the functional group (X) may be introduced in the methacrylic polymer block (a) from the viewpoints of the improvement of heat resistance and thermal decomposition resistance of the acrylic block copolymer (A), and the functional group (X) may be introduced
5 into the acrylic polymer block (b) as reaction points from the viewpoints of imparting oil resistance, further rubber elasticity and compression set durability to the acrylic block copolymer (A). Not particularly limited, it is preferable that either of the block of the
10 methacrylic polymer block (a) or the acrylic polymer block (b) has the functional group (X) from the viewpoints of the control of reaction points, heat resistance, rubber elasticity, mechanical strength, and flexibility.

The content number of the functional group (X) is varied
15 depending on the cohesive force and reactivity of the functional group (X), the structure and composition of the acrylic block copolymer (A), the number of blocks composing the acrylic block copolymer (A), the glass transition temperature, and the site at which the functional group (X) is contained and the mode thereof. Consequently, it may be
20 set according to necessity, and preferably at least 1.0 per one molecule of the acrylic block copolymer (A), and more preferably 2.0 and more. When the number is less than 1.0, the improvement of heat resistance by converting to having a high molecular weight by a bimolecular reaction of the acrylic block copolymer (A) or crosslinking tend to be
25 insufficient.

When the functional group (X) is introduced into the methacrylic polymer block (a), it is preferable to be introduced within a

range in which the moldability of the acrylic block copolymer (A) is not lowered. Since flowage under no pressurization is required particularly in case of the powder slash molding, the cohesive force and the glass transition temperature T_g of the methacrylic polymer block (a) are increased by the introduction of the functional group (X); therefore, melt viscosity is increased and moldability tends to be deteriorated. Specifically, it is preferable to be introduced within a range in which the glass transition temperature T_g of the methacrylic polymer block (a) after introduction of the functional group (X) is preferably at most 130°C , more preferably at most 110°C , and further preferably at most 100°C .

When the functional group (X) is introduced into the acrylic polymer block (b), it is preferably introduced within a range not deteriorating the flexibility, rubber elasticity and low temperature property of the acrylic block copolymer (A). When the cohesive force and the glass transition temperature T_g of the acrylic polymer block (b) are increased by introducing the functional group (X), the flexibility, rubber elasticity and low temperature property tend to be deteriorated. Specifically, it is preferable to be introduced within a range in which the glass transition temperature of the acrylic polymer block (b) after the introduction of the functional group (X) is preferably at most 25°C , more preferably at most 0°C , and further preferably at most -20°C .

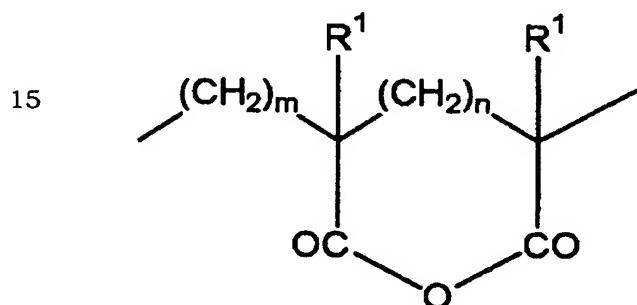
An acid anhydride group, a carboxyl group, a hydroxyl group and an epoxy group, which are preferable as the functional group (X), are explained in the following.

<Acid anhydride group>

Since the acid anhydride group is easily reacted with a

hydroxyl group and an amino group, it can be easily reacted with a compound having those polar groups. Further, when a compound having an active proton is contained in the composition, the acid anhydride group is easily reacted with an epoxy group.

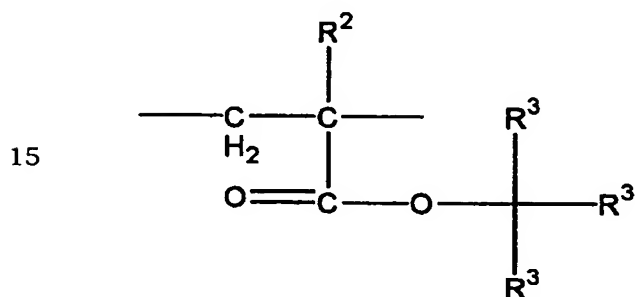
5 The acid anhydride group is not particularly limited, but it may be introduced into the main chain of the acrylic block copolymer (A) and may be introduced into side chains. The acid anhydride group is the acid anhydride group of a carboxyl group, and it is preferably introduced into the main chain from the viewpoint of easiness of
10 introduction into the acrylic block copolymer (A). Specifically, it is represented by the general formula (1). It is more preferable to be contained in a form represented by the general formula (1):



20 (wherein R¹ is hydrogen or a methyl group and may be mutually the same or different. n is an integer of 0 to 3 and m is an integer of 0 or 1). n in the general formula (1) is an integer of 0 to 3, preferably 0 or 1 and more preferably 1. When n is 4 or more, polymerization becomes complicated or cyclization of the acid anhydride group tends
25 to be difficult.

With respect to the introduction method of the acid anhydride group, when a monomer having the acid anhydride group

does not poison a catalyst under polymerization condition, it is preferable to be directly introduced by polymerization, and when a monomer having the acid anhydride group deactivates the catalyst at polymerization, a method of introducing the acid anhydride group by
 5 converting the functional group is preferable. It is not particularly limited, but it is preferable that to be introduced into the acrylic block copolymer in a form which is the precursor of the acid anhydride group, and thereafter cyclized from the viewpoint of easiness of the introduction. It is more preferable to melt-kneading the acrylic block
 10 copolymer which has at least one unit represented by the general formula (2) to be introduced by cyclization;



(wherein R² represents hydrogen or a methyl group. R³ represents
 20 hydrogen, a methyl group or a phenyl group, at least 2 among 3 of R³'s are selected from a methyl group and/or a phenyl group and 3 of R³'s may be mutually the same or different).

The introduction of the unit represented by the general formula (2) to the acrylic block copolymer can be carried out by
 25 copolymerizing an acrylic acid ester or a methacrylic acid ester monomer derived from the general formula (2). Examples of the monomer are t-butyl (meth)acrylate, isopropyl (meth)acrylate,

α,α -dimethylbenzyl (meth)acrylate, α -methylbenzyl (meth)acrylate, but is not limited thereto. Among those, t-butyl (meth)acrylate is preferable from the viewpoints such as availability, easiness of polymerizability, and easiness of generating the acid anhydride group.

5 For the formation of the acid anhydride group, the acrylic block copolymer having the precursor of the acid anhydride group is preferably heated under high temperature and preferably heated at 180 to 300°C. When the temperature is lower than 180°C, generation of the acid anhydride group tends to be insufficient, and when it is
10 higher than 300°C, the acrylic block copolymer in itself having a precursor of the acid anhydride group tends to be decomposed.

<Carboxyl group>

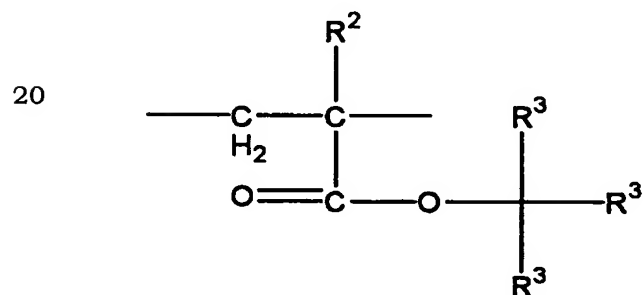
 Since the carboxyl group is easily reacted with an epoxy group and an amino group, it is easily reacted with a compound having
15 those polar groups. The carboxyl group is not particularly limited, but it may be introduced in the main chain of the acrylic block copolymer (A) or may be introduced in a side chain. The carboxyl group is preferably introduced in the main chain from the viewpoint of easiness of the introduction into the acrylic block copolymer (A).

20 Concerning the introduction method of carboxyl group, when a monomer having a carboxyl group does not poison a catalyst under polymerization conditions, the carboxyl group is preferably introduced directly by polymerization, and when a monomer having a carboxyl group deactivates the catalyst at polymerization, a method of
25 introducing the carboxyl group by converting a functional group is preferable.

 In the method of introducing the carboxyl group by

converting a functional group, the carboxyl group is introduced into the acrylic block copolymer in the form of protecting the carboxyl group with an appropriate protecting group or in the form of a functional group being the precursor of the carboxyl group, and then the functional group can be prepared by known chemical reactions. The carboxyl group can be introduced by the method.

As the synthesis method of the acrylic block copolymer (A) having a carboxyl group, for example, there are a method by which an acrylic block copolymer containing a monomer having a functional group such as t-butyl methacrylate, t-butyl acrylate, trimethylsilyl methacrylate or trimethylsilyl acrylate, which is the precursor of the carboxyl group, is synthesized, and the carboxyl group is formed by known chemical reactions such as hydrolysis or acid decomposition (JP-A-10-298248 and JP-A-2001-234146), and a method which a acrylic block copolymer having at least one unit represented by the general formula (2) is melt-kneaded and a carboxyl group is introduced;



(wherein R^2 represents hydrogen or a methyl group. R^3 represents hydrogen, a methyl group or a phenyl group, at least 2 among 3 of R^3 's are selected from a methyl group and/or a phenyl group, and 3 of R^3 's

may be mutually the same or different). The unit represented by the general formula (2) has a route in which a carboxyl group is generated by decomposing the ester unit under a high temperature, and successively, cyclization occurs to generate the acid anhydride group.

- 5 By utilizing this, the carboxyl group can be introduced by appropriately adjusting heating temperature and time in accordance with the kind and content of the unit represented by the general formula (2).

Further, the carboxyl group can be also introduced by hydrolysis of the acid anhydride group.

10 <Hydroxyl group>

- Since a hydroxyl group is easily reacted with an epoxy group, an acid anhydride group and a carboxyl group, it is easily reacted with a compound having those polar groups. The hydroxyl group is not particularly limited, but an alcoholic hydroxyl group is preferable from the viewpoints of the easiness of introduction into the acrylic block copolymer (A), and reactivity.

The hydroxyl group is not particularly limited, but it may be introduced in the main chain of the acrylic block copolymer (A) or may be introduced in the side chain.

- 20 The preferable introduction method of the hydroxyl group is the introduction by directly polymerizing a monomer having a hydroxyl group. Specific examples of the monomer are 2-hydroxyethyl (meth)acrylate, 2-hydroxypropyl (meth)acrylate, 4-hydroxybutyl (meth)acrylate, and the ethylene oxide adduct of (meth)acrylic acid.

25 <Epoxy group>

Since an epoxy group is easily reacted with a carboxyl group, a hydroxyl group and an amino group, it is easily reacted with a

compound having those polar groups. The epoxy group is not particularly limited as long as it is an organic group containing an epoxy ring, specific examples of the epoxy groups are an aliphatic hydrocarbon (for example, alkyl) group having an epoxy ring such as a 1,2-epoxyethyl group, a 2,3-epoxypropyl group (namely, glycidyl group) and a 2,3-epoxy-2-methylpropyl group; and an alicyclic hydrocarbon group having an epoxy ring such as a 3,4-epoxycyclohexyl group. These may be selected according to necessity from the viewpoints of reactivity, reaction speed, and the easiness of availability, and cost.

The epoxy group is not particularly limited, but among those, a glycidyl group is preferable from the viewpoint of easiness of availability.

With respect to the introduction method of an epoxy group, it is preferable to directly introduce a monomer having an epoxy group by polymerization. Examples of the monomer having an epoxy group are the esters of (meth)acrylic acid with an alcohol containing an organic group having an epoxy ring such as glycidyl (meth)acrylate, 2,3-epoxy-2-methylpropyl (meth)acrylate and (3,4-epoxycyclohexyl)methyl (meth)acrylate; and unsaturated compounds containing an epoxy group such as 4-vinyl-1-cyclohexene-1,2-epoxide. These may be selected according to necessity from the viewpoints of reactivity, reaction speed, the easiness of availability, and cost. It is not particularly limited, but among those, glycidyl (meth)acrylate is preferable from the viewpoint of the easiness of availability.

<Methacrylic polymer block (a)>

The methacrylic polymer block (a) is a block obtained by polymerizing a monomer comprising a methacrylic acid ester as the

main component, and preferably comprises 50 to 100 % by weight of the methacrylic acid ester and 0 to 50 % by weight of a vinyl monomer copolymerizable with this. Further, a monomer having the reactive functional group (X) may be contained as the methacrylic acid ester.

- 5 When the proportion of the methacrylic acid ester is less than 50 % by weight, there is a case where weather resistance which are a characteristic of the methacrylic acid ester is deteriorated.

Examples of the methacrylic acid ester composing the methacrylic polymer block (a) are methacrylic acid aliphatic
10 hydrocarbon (for example, alkyl having 1 to 18 carbons) esters such as methyl methacrylate, ethyl methacrylate, n-propyl methacrylate, n-butyl methacrylate, isobutyl methacrylate, n-pentyl methacrylate, n-hexyl methacrylate, n-heptyl methacrylate, n-octyl methacrylate, 2-ethylhexyl methacrylate, nonyl methacrylate, decyl methacrylate,
15 dodecyl methacrylate and stearyl methacrylate; methacrylic acid alicyclic hydrocarbon esters such as cyclohexyl methacrylate and isobornyl methacrylate; methacrylic acid aralkyl esters such as benzyl methacrylate; methacrylic acid aromatic hydrocarbon esters such as phenyl methacrylate and tolyl methacrylate; esters of methacrylic acid
20 with alcohol containing a functional group having ethereal oxygen such as 2-methoxyethyl methacrylate and 3-methoxybutyl methacrylate; methacrylic acid fluorinated alkyl esters such as trifluoromethyl methacrylate, trifluoromethylmethyl methacrylate, 2-trifluoromethylethyl methacrylate, 2-trifluoroethyl methacrylate,
25 2-perfluoroethylethyl methacrylate, 2-perfluoroethyl-2-perfluorobutylethyl methacrylate, 2-perfluoroethyl methacrylate, perfluoromethyl methacrylate, diperfluoromethylmethyl

methacrylate, 2-perfluoromethyl-2-perfluoroethylmethyl methacrylate, 2-perfluorohexylethyl methacrylate, 2-perfluorodecylethyl methacrylate and 2-perfluorohexadecylethyl methacrylate. At least one of these is used. Among those, methyl methacrylate is preferable from the viewpoints of processability, cost, and availability.

Examples of the vinyl monomer copolymerizable with a methacrylic acid ester composing the methacrylic polymer block (a) are an acrylic acid ester, an aromatic alkenyl compound, a vinyl cyanide compound, a conjugated diene compound, a halogen-containing unsaturated compound, an unsaturated carboxylic acid compound, an unsaturated dicarboxylic acid compound, a vinyl ester compound, and a maleimide compound.

Examples of the acrylic acid ester are acrylic acid aliphatic hydrocarbon (for example, alkyl having 1 to 18 carbons) esters such as methyl acrylate, ethyl acrylate, n-propyl acrylate, n-butyl acrylate, isobutyl acrylate, n-pentyl acrylate, n-hexyl acrylate, n-heptyl acrylate, n-octyl acrylate, 2-ethylhexyl acrylate, nonyl acrylate, decyl acrylate, dodecyl acrylate and stearyl acrylate; acrylic acid alicyclic hydrocarbon esters such as cyclohexyl acrylate and isobornyl acrylate; acrylic acid aromatic hydrocarbon esters such as phenyl acrylate and tolyl acrylate; acrylic acid aralkyl esters such as benzyl acrylate; esters of acrylic acid with alcohol containing a functional group having ethereal oxygen such as 2-methoxyethyl acrylate and 3-methoxybutyl acrylate; acrylic acid fluorinated alkyl esters such as trifluoromethylmethyl acrylate, 2-trifluoromethylethyl acrylate, 2-perfluoroethylethyl acrylate, 2-perfluoroethyl-2-perfluorobutylethyl acrylate, 2-perfluoroethyl acrylate, perfluoromethyl acrylate, diperfluoromethylmethyl acrylate,

2-perfluoromethyl-2-perfluoroethylmethyl acrylate,
2-perfluorohexylethyl acrylate, 2-perfluorodecylethyl acrylate and
2-perfluorohexadecylethyl acrylate.

Examples of the aromatic alkenyl compound are styrene,
5 α -methylstyrene, p-methylstyrene, and p-methoxystyrene.

Examples of the vinyl cyanide compound includes
acrylonitrile, and methacrylonitrile.

Examples of the conjugated diene compound are butadiene
and isoprene.

10 Examples of the halogen-containing unsaturated
compound are vinyl chloride, vinylidene chloride, perfluoroethylene,
perfluoropropylene, and vinylidene fluoride.

Examples of the unsaturated carboxylic acid compound
are methacrylic acid and acrylic acid.

15 Examples of the unsaturated dicarboxylic acid compound
are maleic anhydride, maleic acid, a monoalkyl ester and a dialkyl
ester of maleic acid, fumaric acid, and a monoalkyl ester and a dialkyl
ester of fumaric acid.

Examples of the vinyl ester compound are vinyl acetate,
20 vinyl propionate, vinyl pivalate, vinyl benzoate, and vinyl cinnamate.

Examples of the maleimide compound are maleimide,
methylmaleimide, ethylmaleimide, propylmaleimide, butylmaleimide,
hexylmaleimide, octylmaleimide, dodecylmaleimide, stearylmaleimide,
phenylmaleimide, and cyclohexylmaleimide.

25 At least one of these is used. Among those vinyl
monomers, a preferable monomer can be selected from the viewpoints
such as adjustment of a glass transition temperature required for the

methacrylic polymer block (a), and compatibility with the acrylic polymer block (b).

The glass transition temperature of the methacrylic polymer block (a) is preferably 25 to 130°C, more preferably 40 to 110°C, and further preferably 50 to 100°C from the viewpoints of thermal deformation and moldability of the elastomer composition.

From the above-mentioned viewpoints, the methacrylic polymer block (a) is preferably a block obtained by polymerizing methyl methacrylate as the main component, a monomer having the reactive functional group (X) and at least one monomer selected from the group consisting of ethyl acrylate, n-butyl acrylate, and 2-methoxyethyl acrylate for controlling a glass transition temperature.

T_g of the methacrylic polymer block (a) can be set by setting a weight ratio of monomers in a polymer portion according to the above-mentioned Fox formula.

<Acrylic polymer block (b)>

The acrylic polymer block (b) is a block obtained by polymerizing a monomer which comprises an acrylic acid ester as the main component, and preferably comprises 50 to 100 % by weight of an acrylic acid ester and 0 to 50 % by weight of a vinyl monomer copolymerizable with this. Further, a monomer having the reactive functional group (X) may be contained as an acrylic acid ester. When the proportion of the acrylic acid ester is less than 50 % by weight, there may be a case where the physical property of the composition, in particular, flexibility and oil resistance, which are the characteristics of using the acrylic acid ester, is deteriorated.

Examples of an acrylic acid ester composing the acrylic

polymer block (b) are monomers similar to the acrylic acid ester, which is exemplified as a monomer composing the methacrylic polymer block (a).

These can be used alone or in combination of two or more kinds thereof. Among those, n-butyl acrylate is preferable from the viewpoint of the balance of rubber elasticity, property at low temperature and cost. When oil resistance and mechanical property are required, ethyl acrylate is preferable. Further, when imparting low temperature property and oil resistance and improvement of surface tackiness of a resin are required, 2-methoxyethyl acrylate is preferable. Further, when the balance of oil resistance and the low temperature property is required, a combination of ethyl acrylate, n-butyl acrylate, and 2-methoxyethyl acrylate is preferable.

Example of the vinyl monomer copolymerizable with an acrylic acid ester composing the acrylic polymer block (b) are a methacrylic acid ester, an aromatic alkenyl compound, a vinyl cyanide compound, a conjugated diene compound, a halogen-containing unsaturated compound, a silicon-containing unsaturated compound, an unsaturated carboxylic acid compound, an unsaturated dicarboxylic acid compound, a vinyl ester compound, and a maleimide compound. Specific examples thereof are those same as the above-mentioned compounds used for the methacrylic polymer block (a).

At least one of these is used. Among those vinyl monomers, a preferable monomer can be selected from the viewpoint of the balance of glass transition temperature and oil resistance required for the acrylic polymer block (b), and compatibility with the methacrylic

polymer block (a). For example, acrylonitrile can be copolymerized for the purpose of improving oil resistance of the composition.

The glass transition temperature of the acrylic polymer block (b) is preferably at most 25°C, more preferably at most 0°C, and
5 further preferably at most -20°C from the viewpoint of rubber elasticity of the elastomer composition. When the glass transition temperature of the acrylic polymer block (b) is higher than the temperature of environment in which the elastomer composition is used, it is not advantageous because rubber elasticity is hardly expressed.

10 From the above-mentioned viewpoints, the acrylic polymer block (b) is preferably a block obtained by polymerizing a monomer having the reactive functional group (X) and a monomer comprising at least one monomer selected from the group consisting of n-butyl acrylate, ethyl acrylate and 2-methoxyethyl acrylate as the main
15 component.

T_g_b of the acrylic polymer block (b) can be set by setting the weight ratio of each of monomers in a polymer according to the above-mentioned Fox formula.

<Preparation process of acrylic block copolymer (A)>

20 The process for preparing the acrylic block copolymer (A) is not particularly limited, but controlled polymerization using an initiator for a polymer is preferably used. Examples of the controlled polymerization are living anion polymerization, radical polymerization using a chain transfer agent and living radical polymerization which
25 has been recently developed. Among those, the living radical polymerization is preferable from the viewpoint of controlling a molecular weight and a structure of the acrylic block copolymer.

The living radical polymerization is radical polymerization in which activity of the terminal of polymerization is maintained without losing. The living polymerization indicates polymerization in which the terminal always continues to have activity in the narrow
5 sense, but in general, pseudoliving polymerization in which the terminal deactivated and the terminal activated are in equilibrium state is also included. The definition herein is also applied to the latter. Recently, the living radical polymerization has been actively studied by various groups.

10 Example of the living radical polymerization are those using a chain transfer agent such as polysulfide, those using a cobalt porphyrin complex (J. Am. Chem. Soc., 1994, Vol.116, page 7943), those using a radical capturing agent such as a nitroxide compound (Macromolecules, 1994, Vol.27, page 7228), and Atom Transfer Radical
15 Polymerization (ATRP) using a transition metal complex as a catalyst and organic halogenated compound as an initiator. In the present invention, there is no limitation for which method is used, but the Atom Transfer Radical Polymerization is preferable from the viewpoint of easiness of control.

20 The Atom Transfer Radical Polymerization is carried out by using an organic halogenated compound or a halogenated sulfonyl compound as an initiator and a metal complex in which Group VII, Group VIII, Group IX, Group X or Group XI element of the Periodic Table is used as a central metal, as a catalyst (for example, refer to
25 Matyjaszewski et. al., J. Am. Chem. Soc., 1995, Vol.117, page 5614, Macromolecules, 1995, Vol.28, page 7901, Science, 1996, Vol.272, page 866 or Sawamoto et. al., Macromolecules, 1995, Vol.28, page

1721).

According to the method, in general, even though it is radical polymerization in which polymerization speed is very high and a termination reaction such as the coupling of mutual radicals easily occurs, but the polymerization livingly proceeds, a polymer with narrow molecular weight distribution ($M_w/M_n = 1.1$ to 1.5) is obtained and a molecular weight can be freely controlled by changing the ratio of a monomer and an initiator.

In the Atom Transfer Radical Polymerization, as the organic halogenated compound or halogenated sulfonyl compound as an initiator, monofunctional, bifunctional or multifunctional compounds can be used. These may be used in accordance with its purpose, but when a diblock copolymer is produced, a monofunctional compound is preferable from the viewpoint of easy availability of an initiator. When a triblock copolymer of a-b-a type and a triblock copolymer of b-a-b type are produced, a bifunctional compound is preferably used from the viewpoint of reducing the number of reaction steps and time. When a branched block copolymer is produced, a multifunctional compound is preferably used from the viewpoint of reducing the number of reaction steps and time.

Further, a macroinitiator can be also used as the above-mentioned initiator. The macroinitiator is a compound comprising a polymer in which a halogen atom is bonded at the terminal of a molecule, among the organic halogenated compound or the halogenated sulfonyl compound. Since such macroinitiator can be also produced by the controlled polymerization method other than the living radical polymerization process, there is a characteristic that a

block copolymer bonded with a polymer obtained by a different polymerization process is obtained.

Examples of the monofunctional compound are compounds represented by

- 5 $\text{C}_6\text{H}_5\text{-CH}_2\text{X}$,
 $\text{C}_6\text{H}_5\text{-C(H)(X)-CH}_3$,
 $\text{C}_6\text{H}_5\text{-C(X)(CH}_3)_2$,
 $\text{R}^4\text{-C(H)(X)-COOR}^5$,
 $\text{R}^4\text{-C(CH}_3\text{)(X)-COOR}^5$,
10 $\text{R}^4\text{-C(H)(X)-CO-R}^5$,
 $\text{R}^4\text{-C(CH}_3\text{)(X)-CO-R}^5$, and
 $\text{R}^4\text{-C}_6\text{H}_4\text{-SO}_2\text{X}$.

- Wherein, C_6H_5 represents a phenyl group and C_6H_4 represents a phenylene group (it may be either of ortho substitution,
15 meta substitution and para substitution). R^4 represents a hydrogen atom, an alkyl group having 1 to 20 carbons, an aryl group having 6 to 20 carbons or an aralkyl group having 7 to 20 carbons. X represents chlorine, bromine or iodine. R^5 represents a monovalent organic group having 1 to 20 carbons.

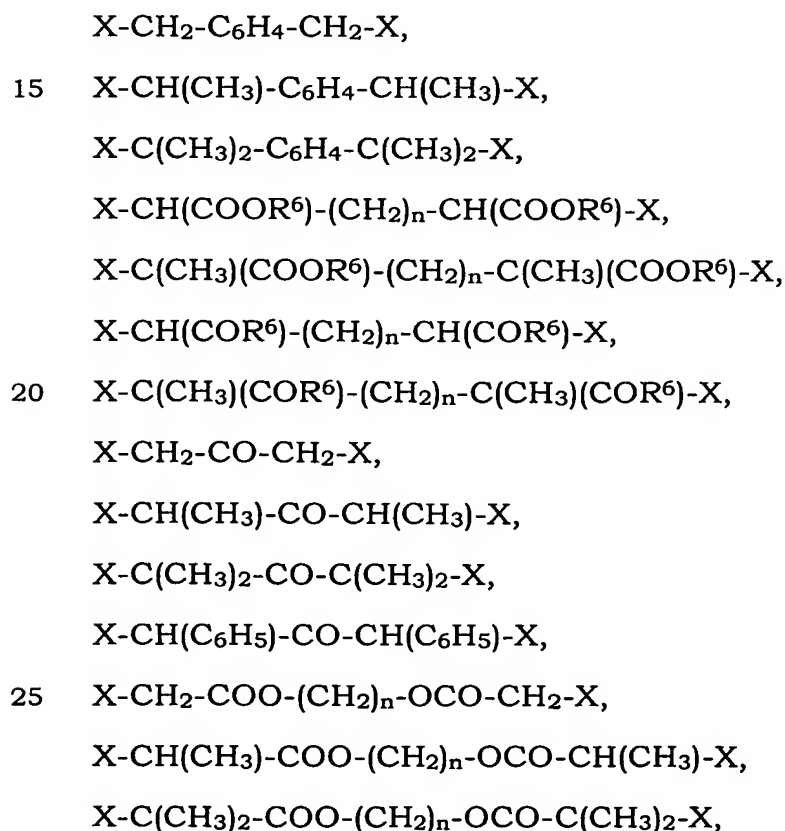
- 20 As R^4 , the specific example of an alkyl group (including a alicyclic hydrocarbon group) having 1 to 20 carbons includes a methyl group, an ethyl group, a propyl group, an isopropyl group, a n-butyl group, an isobutyl group, a tert-butyl group, a n-pentyl group, a n-hexyl group, a cyclohexyl group, a n-heptyl group, a n-octyl group, a
25 2-ethylhexyl group, a nonyl group, a decyl group, a dodecyl group, and an isobornyl group. The specific example of an aryl group having 6 to 20 carbons includes a phenyl group, a tolyl group, and a naphthyl

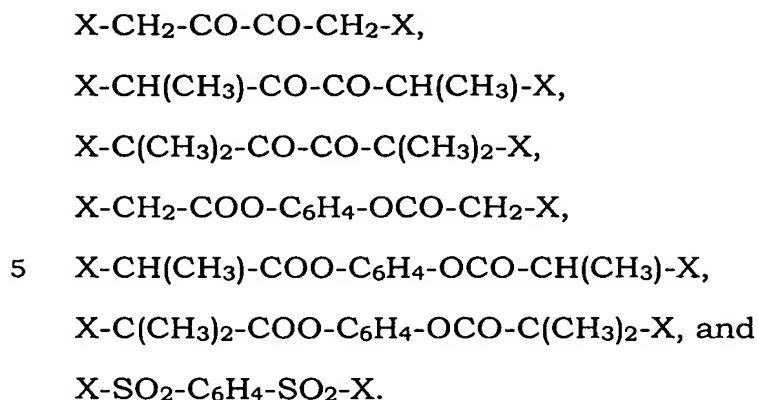
group. Specific examples of an aralkyl group having 7 to 20 carbons are a benzyl group and a phenethyl group.

Specific examples of a monovalent organic group having 1 to 20 carbons being R^5 are groups similar as R^4 .

5 Specific examples of the mono functional compound are tosyl bromide, methyl 2-bromopropionate, ethyl 2-bromopropionate, butyl 2-bromopropionate, methyl 2-bromoisobutyrate, ethyl 2-bromoisobutyrate, and butyl 2-bromoisobutyrate. Among those, ethyl 2-bromopropionate and butyl 2-bromopropionate are preferable
10 from the viewpoint of easiness of controlling polymerization because they are similar as the structure of an acrylic acid ester monomer.

Examples of the bifunctional compound are compounds represented by





Wherein R^6 represents an alkyl group having 1 to 20
 carbons, an aryl group having 6 to 20 carbons or an aralkyl group
 10 having 7 to 20 carbons. The n represents an integer of 0 to 20. C_6H_5 ,
 C_6H_4 and X are similar as the above-description.

Specific examples of an alkyl group having 1 to 20 carbons,
 an aryl group having 6 to 20 carbons and an aralkyl group having 7 to
 20 carbons of R^6 are the same as the specific examples of an alkyl
 15 group having 1 to 20 carbons, an aryl group having 6 to 20 carbons or
 an aralkyl group having 7 to 20 carbons of R^4 .

Specific example of the bifunctional compound are
 bis(bromomethyl)benzene, bis(1-bromoethyl)benzene,
 bis(1-bromoisopropyl)benzene, dimethyl 2,3-dibromosuccinate, diethyl
 20 2,3-dibromosuccinate, dibutyl 2,3-dibromosuccinate, dimethyl
 2,4-dibromoglutarate, diethyl 2,4-dibromoglutarate, dibutyl
 2,4-dibromoglutarate, dimethyl 2,5-dibromoadipate, diethyl
 2,5-dibromoadipate, dibutyl 2,5-dibromoadipate, dimethyl
 2,6-dibromopimelate, diethyl 2,6-dibromopimelate, dibutyl
 25 2,6-dibromopimelate, dimethyl 2,7-dibromosuberate, diethyl
 2,7-dibromosuberate, and dibutyl 2,7-dibromosuberate. Among those,
 bis(bromomethyl)benzene, diethyl 2,5-dibromoadipate and diethyl

2,6-dibromopimelate are preferable from the viewpoint of the availability of raw materials.

Examples of the multifunctional compound are compounds represented by

- 5 $\text{C}_6\text{H}_3-(\text{CH}_2-\text{X})_3$,
 $\text{C}_6\text{H}_3-(\text{CH}(\text{CH}_3)-\text{X})_3$,
 $\text{C}_6\text{H}_3-(\text{C}(\text{CH}_3)_2-\text{X})_3$,
 $\text{C}_6\text{H}_3-(\text{OCO}-\text{CH}_2-\text{X})_3$,
 $\text{C}_6\text{H}_3-(\text{OCO}-\text{CH}(\text{CH}_3)-\text{X})_3$,
10 $\text{C}_6\text{H}_3-(\text{OCO}-\text{C}(\text{CH}_3)_2-\text{X})_3$, and
 $\text{C}_6\text{H}_3-(\text{SO}_2-\text{X})_3$.

Wherein C_6H_3 is a trivalent phenyl group (the position of three bonding hands may be a combination of either of 1-position to 6-position) and X is the same as the above description.

- 15 Specific examples of the multifunctional compound are tris(bromomethyl)benzene and tris(1-bromoethyl)benzene, tris(1-bromoisopropyl)benzene. Among those, tris(bromomethyl)benzene is preferable from the viewpoint of the availability of raw materials.

- 20 Further, when the organic halogenated compound or halogenated sulfonyl compound having a functional group other than a group initiating polymerization is used, a polymer in which a functional group other than a group initiating polymerization is introduced at a terminal or in a molecule can be easily obtained. The
- 25 functional group other than a group initiating polymerization includes an alkenyl group, a hydroxyl group, an epoxy group, an amino group, an amide group, and a silyl group.

As the organic halogenated compound or halogenated sulfonyl compound which can be used as the initiator, carbon atom with which a halogen group (a halogen atom) is bonded with a carbonyl group or a phenyl group and carbon-halogen bond is activated to initiate polymerization. An amount of the initiator used may be determined from the molar ratio with a monomer in conformity with a molecular weight of the acrylic block copolymer required. Namely, the molecular weight of the acrylic block copolymer can be controlled by what molecule of a monomer is used per 1 molecule of the initiator.

The transition metal complex used as the catalyst of the Atom Transfer Radical Polymerization is not particularly limited, but preferable examples are the complex of monovalent or zero-valent copper, the complex of bivalent ruthenium, the complex of bivalent iron and the complex of bivalent nickel.

Among those, the complex of copper is preferable from the viewpoints of cost and reaction control. Examples of the monovalent copper are cuprous chloride, cuprous bromide, cuprous iodide, cuprous cyanide, cuprous oxide, and cuprous perchlorate. Among those, cuprous chloride and cuprous bromide are preferable from the viewpoint of the control of polymerization. When the monovalent copper compound is used, 2,2'-bipyridyl compounds such as 2,2'-bipyridyl and its derivative (such as 4,4'-dinolyl-2,2'-bipyridyl and 4,4'-di(5-nolyl)-2,2'-bipyridyl); 1,10-phenanthroline compounds such as 1,10-phenanthroline and its derivative (such as 4,7-dinolyl-1,10-phenanthroline and 5,6-dinolyl-1,10-phenanthroline); polyamines such as tetramethylethylenediamine (TMEDA), pentamethyldiethylenetriamine and hexamethyl(2-aminoethyl) amine

may be added as a ligand in order to enhance activation of catalyst.

Further, the tris(triphenylphosphin) complex ($\text{RuCl}_2(\text{PPh}_3)_3$) of bivalent ruthenium chloride is also preferable as a catalyst. When the ruthenium compound is used as a catalyst, an aluminum alkoxides may be added as an activating agent. Further, the bistris(triphenylphosphin) complex ($\text{FeCl}_2(\text{PPh}_3)_2$) of bivalent iron, the bistris(triphenylphosphin) complex ($\text{NiCl}_2(\text{PPh}_3)_2$) of bivalent nickel and the bistris(isobutylphosphin) complex ($\text{NiBr}_2(\text{PPh}_3)_2$) of bivalent nickel are also preferable as a catalyst.

The catalyst, ligand and activator used are not particularly limited, but they may be suitably determined by the relation between reaction speed and the initiator, monomer and solvent used. For example, since it is preferable for the polymerization of acrylic monomers such as an acrylic acid ester that the growth terminal of polymer chains has a carbon-bromine bond from the viewpoint of controlling a polymerization, it is preferable that the initiator used is the organic brominated compound or brominated sulfonyl compound and the solvent is acetonitrile, and it is more preferable that a ligand such as pentamethyldiethylenetriamine is used, and a metal complex catalyst in which cupric bromide, preferably copper contained in cuprous bromide is a central metal is used. Further, since it is preferable from the viewpoint of controlling polymerization that the growth terminal of polymer chains has a carbon-chloride bond for the polymerization of methacrylic monomers such as a methacrylic acid ester, the initiator used is the organic chlorinated compound or chlorinated sulfonyl compound, and the solvent is acetonitrile or a mix solvent with toluene if necessary, and it is more preferable that a

ligand such as pentamethyldiethylenetriamine is used and a metal complex catalyst in which cupric chloride, preferably copper contained in cuprous chloride is a central metal is used.

The amounts of the catalyst and ligand used may be
5 determined from the relation between required reaction speed and the amounts of the initiator, monomer and solvent used. For example, when a polymer with a high molecular weight is obtained, the ratio of the initiator to the monomer must be smaller than a case of obtaining a polymer with low molecular weight, but in such a case, the reaction
10 speed can be increased by increasing the catalyst and ligand. Further, when a polymer having a higher glass transition temperature than a room temperature is prepared, an appropriate organic solvent is added for enhancing stirring efficiency by lowering the viscosity of the system. Therefore, the reaction speed tends to be lowered, but in such a case,
15 the reaction speed can be increased by increasing the catalyst and ligand.

The Atom Transfer Radical Polymerization can be carried out in non-solvent (bulk polymerization) or in various solvents. Further, when polymerization is carried out by bulk polymerization or
20 in various solvents, the polymerization can be also stopped on the way.

As the solvent, for example, a hydrocarbon solvent, an ether solvent, a halogenated hydrocarbon solvent, a ketone solvent, an alcohol solvent, a nitrile solvent, an ester solvent, and a carbonate solvent can be used.

25 The hydrocarbon solvent includes benzene and toluene. The ether solvent includes diethyl ether and tetrahydrofuran. The halogenated hydrocarbon solvent includes methylene chloride and

chloroform. The ketone solvent includes acetone, methyl ethyl ketone, and methyl isobutyl ketone. The alcohol solvent includes methanol, ethanol, propanol, isopropanol, and n-butanol and tert-butanol. The nitrile solvent includes acetonitrile, propionitrile, and benzonitrile.

- 5 The ester solvent includes ethyl acetate and butyl acetate. The carbonate solvent includes ethylene carbonate and propylene carbonate.

One or more of the solvents mentioned above may be used.

- When the solvent is used, the amount used may be
10 suitably determined from the relation between the viscosity of the whole system and stirring efficiency required. Further, when polymerization is carried out by bulk polymerization or in various solvents, the conversion of a monomer at stopping the reaction may be suitably determined from the relation between the viscosity of the
15 whole system and required stirring efficiency, even if the polymerization is stopped on the way.

The polymerization can be carried out in a range of 23°C to 200°C and preferably in a range of 50°C to 150°C.

- In order to produce the acrylic block copolymer by the
20 polymerization, there are mentioned a method of successively adding a monomer, a method of polymerizing the next block using a polymer preliminarily synthesized, as a macroinitiator, a method of bonding polymers separately polymerized by a reaction. Either of these methods may be used and may be used in accordance with purposes.
25 The method of successively adding a monomer is preferable from the viewpoint of simplicity and easiness of the preparation steps.

The reaction solution obtained by the polymerization

contains a mixture of a polymer and the metal complex. An organic acid containing a carboxyl group or a sulfonyl group is added thereto to prepare a salt with the metal complex, and the solid of the salt with the metal complex prepared is removed by filtration, successively, 5 impurities such as an acid remaining in the solution are removed by basic active alumina, a basic absorbent, a solid inorganic acid, an anion exchange resin and cellulose anion exchanger adsorption treatment and thereby, the solution of the acrylic block copolymer can be obtained.

10 The polymerization solvent and unreacted monomer are successively removed by evaporation from the polymer solution obtained in this manner, and the acrylic block copolymer is isolated. As the evaporation system, a thin film evaporation system, a flash evaporation system, a horizontal type evaporation system equipped 15 with an extrusion screw can be used. Since the acrylic block copolymer has adhering property, efficient evaporation can be carried out by the horizontal type evaporation system equipped with an extrusion screw alone or a combination with other evaporation system among the above-mentioned evaporation systems.

20 <Compound (B)>

 The compound (B) composing the thermoplastic elastomer composition relating to the present invention is not particularly limited, it is a compound containing 1.1 or more of the functional groups (Y) in one molecule, and preferably a compound which reacts with the 25 functional group (X), and can convert the acrylic block copolymer (A) to having a high molecular weight by an inter-bimolecular reaction or crosslinking. The number mentioned here represents the average

number of the functional group (Y) existing in the whole compound (B).

In the present invention, the functional group (Y) is not particularly limited so far as it is a group reacting with the functional group (X) in the block copolymer (A), and at least one kind selected
5 from an epoxy group, a carboxyl group, a hydroxyl group, an amino group and an acid anhydride group is preferable from the viewpoints of stability of a bond generated by the reaction, easiness of the reaction, and the cost of the compound (B).

Two or more kinds of these functional groups (Y) can be
10 also used in combination, but when two or more kinds are used in combination, it is preferable that they do not disturb the reaction with the functional group (X) by a mutual reaction.

The content number of the functional group (Y) in the compound (B) is changed depending on the reactivity of the functional
15 group (Y), a site in which the functional group (Y) is contained and its mode. Accordingly, it may be set according to requirements, but is preferably 1.1 or more in the compounds (B), more preferably 1.5 or more, and particularly preferably 2.0 or more. When it is less than 1.1, effect as a high molecular weight-converting reaction agent for the
20 inter-bimolecular reaction of the acrylic block copolymer or as a crosslinking agent is low and the improvement of heat resistance of the acrylic block copolymer (A) tends to be insufficient.

When the functional group (Y) is an epoxy group, the compound (B) having an epoxy group is not particularly limited so far
25 as it is a compound containing 1.1 or more of epoxy groups in one molecule. There are exemplified epoxy resins such as a bisphenol A type epoxy resin, a bisphenol F type epoxy resin, a bisphenol AD type

epoxy resin, a bisphenol S type epoxy resin, a hydrogenated epoxy resin thereof, a glycidyl ester type epoxy resin, a glycidylamine type epoxy resin, an alicyclic type epoxy resin, a novolak type epoxy resin, a urethane modified epoxy resin having a urethane bond, a fluorinated epoxy resin, a rubber modified epoxy resin containing a polybutadiene or NBR, a flame retardant type epoxy resin such as glycidyl ether of tetrabromobisphenol A; epoxy plasticizers such as epoxidized soy bean oil, epoxidized linseed oil and an epoxidized fatty acid alkyl ester which are the glycidyl ethers of polyvalent alcohol or the glycidyl esters of polybasic acid; polymers containing an epoxy group such as BONDFAST (trade name, available from Sumitomo Chemical Co., Ltd.); the polymers containing an epoxy group of unsaturated polymers such as an olefin polymer, a styrene polymer and an acrylic polymer. However, it is not limited thereto and compounds containing an epoxy group which are generally used can be used. These compounds containing an epoxy group may be used alone or at least two kinds thereof may be used in combination.

When the functional groups (Y) is a carboxyl group, the compound (B) having carboxyl groups is not particularly limited so far as it is a compound containing 1.1 or more of carboxyl groups in a molecule. Examples are polymers containing a carboxyl group such as adipic acid, itaconic acid, iminodiacetic acid, glutaric acid, succinic acid, citraconic acid, oxalic acid, tartaric acid, sebacic acid, azelaic acid, decanedicarboxylic acid, dodecanedicarboxylic acid, cyclohexanedicarboxylic acid, trans-1,2-cyclohexanediamine tetraacetic acid, fumaric acid, brassilic acid (n11), malonic acid (n1), citraconic acid, maleic acid, isophthalic acid, terephthalic acid,

1,4-naphthalene dicarboxylic acid, 2,6-naphthalene dicarboxylic acid,
5-hydroxyisophthalic acid, o-phthalic acid, 3,3',4,4'-benzophenone
tetracarboxylic acid, diphenoxyethane dicarboxylic acid, biphenyl
dicarboxylic acid, diphenyl ether dicarboxylic acid, diphenylsulfone
5 dicarboxylic acid, diphenylketone dicarboxylic acid, anthracene
dicarboxylic acid, α,β -bis(4-carboxyphenoxy)ethane, trimellitic acid,
pyromellitic acid, imidazole-4,5-dicarboxylic acid, chelidamic acid,
2,3-pyrazine dicarboxylic acid, citric acid, glycyrrhizic acid, asparagic
acid, glutamic acid, malic acid, polymers containing a carboxyl group
10 such as AQUPEC (trade name, available from Sumitomo Seika
Chemicals Co., Ltd.) and ACTFLOW (trade name, available from Soken
Chemical & Engineering Co., Ltd.); polymers containing a carboxyl
group of unsaturated polymer such as petroleum resins such as an
olefin polymer, a styrene polymer and an acrylic polymer. However, it
15 is not particularly limited thereto and compounds containing a
carboxyl group which are generally used can be used. These
compounds containing carboxyl groups may be used alone or at least
two kinds thereof may be used in combination.

When the functional groups (Y) is an amino group, the
20 compound (B) having amino groups is not particularly limited so far as
it is a compound containing at least 1.1 or more of amino groups in a
molecule. Examples are aliphatic polyamines such as
ethylenediamine, diethylenetriamine, triethylenetetramine,
tetraethylenepentamine, pentaethylenehexamine,
25 tetramethylenediamine, pentamethylenediamine,
hexamethylenediamine, trimethylehexamethylenediamine,
dodecamethylenediamine, polyoxypropylenepolyamine,

1,2-diaminopropane, bis(hexamethylene)triamine,
tris(2-aminoethyl)amine, N,N'-dimethylethylenediamine,
N,N'-diethyl-1,3-propanediamine, N,N'-dibutyl-1,6-hexanediamine,
N,N'-diethyl-2-buten-1,4-diamine, N,N'-diethyl-1,4-diaminocyclohexane
5 and polyether polyamine with a number average molecular weight of
200 to 1000 (such as polyoxypropylenediamine having amino groups at
both terminals); alicyclic polyamines such as cyclohexylenediamine,
piperazine, isophoronediamine, 1,3-bisaminomethylcyclohexane,
1,4-bisaminopropylpyperazine, 1,3-bisaminocyclohexane,
10 di(aminodicyclohexyl)methane, ,
3,3'-dimethyldi(aminocyclohexyl)methane,
1-cyclohexylamino-3-aminopropane, 1,4-diaminocyclohexane,
1,4-diaminocyclopentane; di(aminocyclohexyl)methane,
di(aminocyclohexyl)sulfone, 1,3-di(aminocyclohexyl)propane,
15 4-isopropyl-1,2-diaminocyclohexane, 2,4-diaminocyclohexane,
N,N'-diethyl-1,4-diaminocyclohexane,
3,3'-dimethyl-4,4'-diaminodicyclohexylmethane, and
3-aminomethyl-3,3,5-trimethylcyclohexylamine; aromatic polyamines
such as melamine, benzidine, o-phenylenediamine,
20 m-phenylenediamine, p-phenylenediamine,
4,4'-diaminodiphenylmethane, 4,4'-diaminodiphenylsulfone,
4,4'-diaminodiphenylether, 2,4-diaminodiphenylamine,
1,5-diaminonaphthalene, 1,8-diaminonaphthalene and
2,4-diaminotoluene; the polymers containing an amino group of
25 unsaturated polymers such as petroleum resins such as an olefin
polymer, a styrene polymer and an acrylic polymer. However, it is not
limited thereto. These compounds containing amino groups may be

used alone or at least two kinds thereof may be used in combination.

When the functional groups (Y) is a hydroxyl group, the compound (B) having hydroxyl groups is not particularly limited so far as it is a compound containing at least 1.1 or more of hydroxyl groups in a molecule. Examples are aliphatic compounds containing bivalent hydroxyl group such as ethylene glycol, 1,2-propanediol, 1,3-propanediol, 1,2-butanediol, 1,3-butanediol, 1,4-butanediol, 2,3-butanediol, 2,4-pentanediol, 1,5-pentanediol, 1,4-butanediol, 1,6-hexanediol, 2-butyl-2-ethyl-1,3-propanediol, propylene glycol, dipropylene glycol, tripropylene glycol, 1,6-hexanediol, diethylene glycol, dipropylene glycol, trimethylene glycol, triethylene glycol, tetraethylene glycol, 3,4-hexylene glycol, 1,3-butylene glycol, octanediol, 3,6-dimethyl-4-octene-3,6-diol, 2,5-dimethyl-3-hexyne-2,5-diol, 2,5-dimethyl-2,5-hexanediol, isoprene glycol, 3-chloro-1,2-propanediol, 1,4-cyclohexanediol, cyclopentanol, 1,3-dihydroxyacetone, disodium 1,4-dihydroxy-1,4-butanedisulfonate, thioglycol, neopentylglycol, hexylene glycol, polyethylene glycol, polytetramethylene ether glycol, polypropylene glycol, 3-methyl-1,5-pentanediol, glycerine monoacetate, glycerine monobutylate, glycerine- α -monomethyl ether, and glycerine- α -monobutyl ether, aromatic compounds containing bivalent hydroxyl group such as catechol, 1,4-dihydroxyanthraquinone, hydroquinone, 1,4-dihydroxynaphthalene, bisphenol A, bisphenol S, 2,3,5-trimethylhydroquinone, p-hydroxyphenethyl alcohol, protocatechuic acid, and resorcinol, polymer containing bivalent hydroxyl group such as polyethylene oxide, aliphatic compounds containing trivalent hydroxyl group such as trimethylolethane, trimethylolpropane, 1,2,6-hexanetriol, and glycerine, aromatic

compounds containing trivalent hydroxyl group such as 2,3,4-trihydroxybenzophenone, phloroglucinol, and lauryl gallate, aliphatic compounds containing tetravalent hydroxyl group such as pentaerythritol, diglycerine, compounds containing polyvalent hydroxyl
5 group such as sorbitol, polyglycerine, polyvinylalcohol, ethylene-vinylalcohol copolymer, polyparaviny phenol, and polyvinyl butylal, polymer containing hydroxyl group such as ACTFLOW (trade name, available from Soken Chemical & Engineering Co., Ltd.); the polymers containing a hydroxyl group of unsaturated polymer such as
10 petroleum resins such as an olefin polymer, a styrene polymer and an acrylic polymer. However, it is not particularly limited to these. These compounds containing hydroxyl groups may be used alone or two or more kinds thereof may be used in combination.

When the functional groups (Y) is an acid anhydride group,
15 the compound (B) having acid anhydride groups is not particularly limited so far as it is a compound containing 1.1 or more of acid anhydride groups in a molecule. Examples are 3,3',4,4'-biphenyltetracarboxylic acid dianhydride, 3,3',4,4'-benzophenonetetracarboxylic acid dianhydride, pyromellitic
20 acid dianhydride, 3,3',4,4'-diphenylsulfonetetracarboxylic acid dianhydride, maleinated methylcyclohexane tetrabasic acid anhydride, polymers containing acid anhydride groups such as isobutylene maleic anhydride copolymer and BONDINE (trade name, available from Sumika ATOFINA Inc.), but it is not limited thereto. These
25 compounds containing acid anhydride groups may be used alone or two or more kinds may be used in combination.

When the functional groups (Y) is an oxazoline group, the

compound (B) having oxazoline groups is not particularly limited so far as it is a compound containing 1.1 or more of oxazoline groups in a molecule. Examples are multifunctional oxazoline compounds such as 2,2'-bis(2-oxazoline), 2,2'-methylene-bis(2-oxazoline),
5 2,2'-ethylene-bis(2-oxazoline), 2,2'-trimethylene-bis(2-oxazoline), 2,2'-tetramethylene-bis(2-oxazoline), 2,2'-hexamethylene-bis(2-oxazoline), 2,2'-octamethylene-bis(2-oxazoline), 2,2'-ethylene-bis(4,4'-dimethyl-2-oxazoline),
10 2,2'-p-phenylene-bis(2-oxazoline), 2,2'-m-phenylene-bis(2-oxazoline), 2,2'-m-phenylene-bis(4,4'-dimethyl-2-oxazoline), 1,3-phenylene-bisoxazoline, 1,4-phenylene-bisoxazoline, bis(2-oxazolinylcyclohexane)sulfide and bis(2-oxazolinylnorbornane)sulfide; polymers containing oxazoline
15 groups such as EPOCROS (trade name, available from Nippon Shokubai Co., Ltd.), but it is not limited to these. These compounds containing oxazoline groups may be used alone or two or more kinds thereof may be used in combination.

The compound (B) of the present invention is not
20 particularly limited, but it is preferably a compound which improves molding flowability as a plasticizer at the molding of the composition and can simultaneously react with the acrylic block copolymer (A).

Further, the compound (B) of the present invention is preferably a compound having a boiling point of at least 200°C, more
25 preferably at least 230°C, and further preferably at least 250°C. Since the obtained composition is molded at a high temperature, the compound (B) is easily evaporated at molding when the boiling point is

at most 200°C, and a molding method and conditions thereof are limited.

Further, the compound (B) of the present invention is preferably a polymer and is preferably a polymer having a weight
5 average molecular weight of 50,000 or less. When the weight average molecular weight is more than 50,000, the effect of improving the molding flowability as a plasticizer tends to be lowered.

Among the above-mentioned compound having an epoxy group, preferable examples are epoxy plasticizers such as a bisphenol
10 A type epoxy resin, an epoxidized soy bean oil, an epoxidized linseed oil and an epoxidized fatty acid alkyl ester which are the glycidyl ethers of a polyvalent alcohol or the glycidyl esters of polybasic acid, from the viewpoints of compatibility with the acrylic block copolymer (A), the easiness of availability, cost, low volatility at molding, the improvement
15 effect of moldability, and the mechanical strength of the obtained molded article. Further, among the above-mentioned compound having carboxylic groups, the preferable example includes a trimellitic acid and polymers containing a carboxylic group such as ACTFLOW (trade name, available from Soken Chemical & Engineering Co., Ltd.).
20 Further, among the above-mentioned compound having amino groups, preferable examples are hexamethylenediamine and polyether polyamine with a number average molecular weight of 200 to 1000. Further, among the above-mentioned compound having hydroxyl groups, preferable examples are polyethylene glycol, polypropylene
25 glycol, glycerin and polymers containing a hydroxyl group such as ACTFLOW (trade name, available from Soken Chemical & Engineering Co., Ltd.). Further, among the above-mentioned compound having

acid anhydride groups, preferable examples are 3,3',4,4'-biphenyltetracarboxylic acid dianhydride and pyromellitic acid dianhydride. Further, among the above-mentioned compound having oxazoline groups, a preferable example is 1,3-phenylene-bisoxazoline.

5 The compound (B) is preferably used in a range of 0.1 to 100 parts by weight based on 100 parts by weight of the acrylic block copolymer (A), more preferably a range of 1 to 50 parts by weight, and particularly preferably a range of 1.5 to 20 parts by weight. When the compounding amount is less than 0.1 parts by weight, moldability and
10 the heat resistance of the obtained molded article are occasionally insufficient, and when it exceeds 100 parts by weight, the mechanical property of the obtained composition tends to be lowered.

<Thermoplastic elastomer composition>

 The thermoplastic elastomer composition of the present
15 invention is the thermoplastic elastomer composition comprising the acrylic block copolymer (A) having a functional group (X) and the compound (B) containing 1.1 or more of functional groups (Y) in one molecule. The functional group (X) is preferably at least one kind of functional groups selected from an acid anhydride group, a carboxyl
20 group, a hydroxyl group and an epoxy group. The functional group (Y) is preferably at least one kind of functional groups selected from an epoxy group, a carboxyl group, a hydroxyl group, an amino group, an acid anhydride group and an oxazoline group.

 Not particularly limited, but among those, a combination
25 that the functional group (X) is at least one kind of functional groups selected from an acid anhydride group and a carboxyl group and the functional group (Y) is an epoxy group is preferable from the viewpoints

of the stability of a bond formed by the reaction, easiness of the reaction, cost, moldability, and heat resistance and thermal stability of the obtained molded article.

The thermoplastic elastomer composition of the present invention is low in melt viscosity at molding and excellent in moldability, and on the other hand, the functional group (X) and the functional group (Y) are reacted at molding and the acrylic block copolymer (A) is preferably converted to having a high molecular weight or is crosslinked. It is more preferable to be crosslinked at molding from the viewpoint of the improvement of heat resistance.

In the thermoplastic elastomer composition of the present invention, various additives and catalysts may be added if necessary in order to accelerate a reaction at molding. For example, when the functional group (X) or the functional group (Y) is an epoxy group, curing agents such as acid anhydride, amine and imidazole which are generally used for an epoxy resin can be also used, and when the functional group (X) or the functional group (Y) is a hydroxyl group or a carboxyl group, known esterification catalysts and known ester exchange catalysts such as bivalent tin compounds and titanate acid esters can be used.

In the thermoplastic elastomer composition of the present invention, a filler is compounded if necessary, and it can be preferably used. The filler is not particularly limited, and examples are reinforcing fillers such as wood flour, pulp, cotton chip, asbestos, glass fiber, carbon fiber, mica, walnut shell flour, rice hull flour, graphite, diatom earth, white clay, silica (fumed silica, precipitating silica, crystal silica, fused silica, dolomite, silicic anhydride, hydrated silicic

acid) and carbon black; fillers such as heavy calcium carbonate, colloidal calcium carbonate, magnesium carbonate, diatom earth, calcined clay, clay, talc, titanium oxide, bentonite, organic bentonite, ferric oxide, colcothar, aluminum fine powder, flint powder, zinc oxide,
5 active zinc oxide, zinc powder, zinc carbonate and sand bur balloon; fibrous fillers such as asbestos, glass fiber and glass filament, carbon fiber, Kevlar fiber and polyethylene fiber,.

Among those fillers, inorganic fillers are more preferable from the viewpoints of the improvement of mechanical property, reinforcing effect, cost, and titanium oxide, carbon black, calcium
10 carbonate, silica and talc are further preferable.

Further, in case of silica, silica on which the surface is preliminarily hydrophobically treated with organosilicone compounds such as organosilanes, organosilazane and diorganopolysiloxane may
15 be used. Further, as calcium carbonate, there may be also used calcium carbonate in which surface treatment is carried out using surface treating agents such as organic substances such as fatty acids, fatty acid soap and fatty acid esters; various surfactants, and various coupling agents such as a silane coupling agent and a titanate
20 coupling agent.

An amount to be added in the case of using the filler is preferably in a range of 5 to 200 parts by weight based on 100 parts by weight of the acrylic block copolymer (A), and more preferably in a range of 10 to 100 parts by weight. When the amount is less than 5
25 parts by weight, the reinforcing effect of the obtained molded article is occasionally not sufficient, and when it exceeds 200 parts by weight, moldability of the composition tends to be lowered. At least one filler

can be used.

The thermoplastic elastomer composition of the present invention can compound various lubricants for moldability, mold releasing property and for making the surface of a molded article low frictional, and it can be preferably used.

The lubricant includes fatty acids such as stearic acid and palmitic acid; metal salts of fatty acids such as calcium stearate, zinc stearate, magnesium stearate, potassium palmitate and sodium palmitate; waxes such as polyethylene wax, polypropylene wax and montanic acid wax; low molecular weight polyolefins such as low molecular weight polyethylene and low molecular weight polypropylene; polyorganosioxanes such as dimethylpolysiloxane; cucutadecylamine, alkyl phosphate, fatty acid esters, amide lubricants such as ethylenebisstearylamine, fluorine resin powder such as ethylene tetrafluoride resin, molybdenum disulfide powder, silicone resin powder, silicone rubber powder, and silica. One or more kind of these can be used. Among those, stearic acid, zinc stearate, calcium stearate, fatty acid esters, ethylenebisstearylamine which are excellent in cost and moldability are preferable.

An amount to be added in the case of using the lubricant is preferably in a range of 0.1 to 20 parts by weight based on 100 parts by weight of the acrylic block copolymer (A), and more preferably in a range of 0.2 to 20 parts by weight. When the compounding amount is less than 0.1 part by weight, the improving effect of moldability and making the obtained molded article have low friction are occasionally not sufficient, and when it exceeds 20 parts by weight, the mechanical property and chemical resistance of the obtained molded article tend to

be deteriorated. One or more lubricant can be used.

In the thermoplastic elastomer composition of the present invention, various additives other than those described above may be added if necessary, for the purpose of the adjustment of various physical properties of the thermoplastic elastomer composition and the obtained molded article. As the additives, a stabilizer, a plasticizer, a flexibility imparting agent, a flame retardant, a pigment, an antistatic agent, an antibacterial and antifungus agent may be added.

As the above-mentioned stabilizer, an antioxidant, a photo stabilizer, and an ultraviolet absorbent are raised. Examples of antioxidant are amine antioxidants such as phenyl- α -naphthylamine (PAN), octyldiphenylamine, N,N'-diphenyl-p-phenylenediamine (DPPD), N,N'-di- β -naphthyl-p-phenylenediamine (DNPD), N-(1,3-dimethylbutyl)-N'-phenyl-p-phenylenediamine, N-phenyl-N'-isopropyl-p-phenylenediamine (IPPN), N,N'-diallyl-p-phenylenediamine, phenothiazine derivative, diallyl-p-phenylenediamine mixture, alkylated phenylenediamine, 4,4'-bis(α,α -dimethylbenzyl)diphenylamine, N-phenyl-N'-(3-methacryloyloxy-2-hydropropyl)-p-phenylenediamine, diallylphenylenediamine mixture, diallyl-p-phenylenediamine mixture, N-(1-methylheptyl)-N'-phenyl-p-phenylenediamine and diphenylamine derivative; imidazole antioxidants such as 2-mercaptobenzoimidazole (MBI); phenol antioxidants such as 2,6-di-t-butyl-4-methylphenol and pentaerythrityltetrakis[3-(5-di-t-butyl-4-hydroxyphenol)-propionate]; phosphate antioxidants such as nickel diethyl-dithiocarbamate; the secondary antioxidants such as triphenylphosphite; 2-t-butyl-6-(3-t-butyl-2-hydroxy-5-methylbenzyl)-4-methylphenyl

acrylate, and
2-[1-(2-hydroxy-3,5-di-t-pentylphenyl)ethyl]-4,6-di-t-pentylphenyl
acrylate. Further, examples of the photo stabilizer and ultraviolet
absorbent are 4-t-butylphenyl salicylate, 2,4-dihydroxybenzophenone,
5 2,2'-dihydroxy-4-methoxybenzophenone, ethyl-2-cyano-3,3'-diphenyl
acrylate, 2-ethylhexyl-2-cyano-3,3'-diphenyl acrylate,
2-hydroxy-5-chlorobenzophenone,
2-hydroxy-4-methoxybenzophenone-2-hydroxy-4-octoxybenzophenone,
monoglycol salicylate, oxalic amide, and
10 2,2',4,4'-tetrahydroxybenzophenone.

Examples of industrial products are Irganox 1010
(available from Chiba Specialty Chemicals Co., Ltd.), SANOL LS770
(available from SANKYO Lifetech Co., Ltd.), ADEKA STAB LA-57
(available from ADEKA Corporation), ADEKA STAB LA-68 (available
15 from ADEKA Corporation), Chimassorb 944 (available from Chiba
Specialty Chemicals Co., Ltd.), SANOL LS765 (available from SANKYO
Lifetech Co., Ltd.), ADEKA STAB LA-62 (available from ADEKA
Corporation), TINUVIN 144 (available from Chiba Specialty Chemicals
Co., Ltd.), ADEKA STAB LA-63 (available from ADEKA Corporation),
20 TINUVIN 622 (available from Chiba Specialty Chemicals Co., Ltd.),
ADEKASTAB LA-32 (available from ADEKA Corporation), ADEKA STAB
LA-36 (available from ADEKA Corporation), TINUVIN 571 (available
from Chiba Specialty Chemicals Co., Ltd.), TINUVIN 234 (available from
Chiba Specialty Chemicals Co., Ltd.), ADEKA STAB LA-31 (available
25 from ADEKA Corporation.), TINUVIN 1130 (available from Chiba
Specialty Chemicals Co., Ltd.), ADEKA STAB AO-20 (available from
ADEKA Corporation.), ADEKA STAB AO-50 (available from ADEKA

Corporation.), ADEKA STAB 2112 (available from ADEKA Corporation.),
ADEKA STAB PEP-36 (available from ADEKA Corporation.),
SUMILIZER GM (available from Sumitomo Chemical Co., Ltd.),
SUMILIZER GS (available from Sumitomo Chemical Co., Ltd.), and
5 SUMILIZER TP-D (available from Sumitomo Chemical Co., Ltd.).
These may be used alone and at least two kinds thereof may be used in
combination. Among those, SANOL LS770, Irganox 1010, SUMILIZER
GS and TINUVIN 234 are preferable from the viewpoints of the
prevention effect of deterioration by heat and light of the acrylic block
10 and the cost thereof.

Example of the above-mentioned plasticizer are phthalic
acid derivatives such as dimethyl phthalate, diethyl phthalate,
di-n-butyl phthalate, di-(2-ethylhexyl) phthalate, diheptyl phthalate,
diisodecyl phthalate, di-n-octyl phthalate, diisononyl phthalate,
15 ditridecyl phthalate, octyldecyl phthalate, butylbenzyl phthalate and
dicyclohexyl phthalate; isophthalic acid derivatives such as dimethyl
isophthalate; tetrahydrophthalic acid derivatives such as
di-(2-ethylhexyl) tetrahydrophthalate; adipic acid derivatives such as
dimethyl adipate, dibutyl adipate, di-n-hexyl adipate, di-(2-ethylhexyl)
20 adipate, isononyl adipate, diisodecyl adipate and dibutyldiglycol
adipate; azelaic acid derivatives such as di-(2-ethylhexyl) azelate;
sebacic acid derivatives such as dibutyl sebacate; dodecan-2-ic acid
derivatives; maleic acid derivatives such as dibutyl maleate and
di-(2-ethylhexyl) maleate; fumaric acid derivatives such as dibutyl
25 fumarate; trimellitic acid derivatives such as tris-(2-ethylhexyl)
trimellitate; pyromellitic acid derivatives; citric acid derivatives such as
acetyltributyl citrate; itaconic acid derivatives; oleic acid derivatives;

ricinoleic acid derivatives; stearic acid derivatives; other fatty acid derivatives; sulfonic acid derivatives; phosphoric acid derivatives; glutaric acid derivatives; polyester plasticizers which are polymers of dibasic acids such as adipic acid, azelaic acid and phthalic acid with glycol and monovalent alcohol; glycol derivatives; glycerin derivatives; paraffin derivatives such as chlorinated paraffin; epoxy derivative polyester polymerization type plasticizers; polyether polymerization type plasticizers; carbonate derivatives such as ethylene carbonate and propylene carbonate. The plasticizer in the present invention is not limited thereto, various plasticizers can be used and those which are widely commercially available as a plasticizer for rubber can be also used. It can be expected that these compounds can lower the viscosity of the acrylic block copolymer (A). Examples of commercially available plasticizers are THIOCOL TP (available from Morton Co.), ADEKACIZER O-130P, C-79, UL-100, P-200, and RS-735 (available from ADEKA Corporation.). Examples of a high molecular weight plasticizer other than those are acrylic polymer, polypropylene glycol polymer, polytetrahydrofuran polymer, and polyisobutylene polymer. It is not particularly limited, but among those, adipic acid derivatives, phthalic acid derivatives, glutaric acid derivatives, trimellitic acid derivatives, pyromellitic acid derivatives, polyether plasticizers, glycerin derivatives, epoxy derivative polyester polymerization type plasticizers, polyether polymerization type plasticizers are preferable since they have low volatility and weight loss thereof by heating is small.

The above-mentioned flexibility imparting agent is not particularly limited, and examples are a softening agent such as process oil; oil such as animal oil and plant oil; petroleum fractions

such as kerosene, light oil, heavy oil and naphtha. As the softening agent, process oil is mentioned, and more specific examples are paraffin oil; naphthene process oil; and petroleum process oil such as aromatic process oil. As the plant oil, examples are ricinus oil, cotton
5 seed oil, faxseed oil, rape seed oil, soy bean oil, palm oil, coconut oil, peanut oil, pine oil, and tall oil, and one or more of these flexibility imparting agents can be used.

As the above-mentioned flame retardant, the following compounds can be mentioned but they are not particularly limited,
10 and examples are triphenyl phosphate, tricresyl phosphate, decabromobiphenyl, decabromobiphenyl ether, and antimony trioxide. These can be used alone, or a plurality of these may be used in combination.

As the above-mentioned pigment, the following compounds
15 can be mentioned but they are not particularly limited, and examples are carbon black, titanium oxide, zinc sulfide, and zinc oxide. These can be used alone, or a plurality of these may be used in combination.

<Preparation process of thermoplastic elastomer composition>

The preparation process of thermoplastic elastomer
20 composition is not particularly limited, but for example, as a batch type kneading device, a mixing roll, a Banbury mixer, a pressuring kneader and a high shearing mixer can be used, and as a continuous kneading equipment, a uniaxial extruder, a biaxial extruder, and a KCK extruder may be used. Further, known processes such as a
25 process of mechanically mixing a composition to be formed in pellets can be used.

A temperature at kneading for preparing the thermoplastic

elastomer composition is preferably a temperature at which the acrylic block copolymer (A) is reacted with the compound (B) and moldability is not lowered. A temperature at which the acrylic block copolymer (A) is reacted with the compound (B) and moldability is deteriorated is
5 determined by kinds of the functional group (X) and the functional group (Y), introduction amount, the composition of the acrylic block copolymer (A) and the compound (B), and the compatibility of the acrylic block copolymer (A) with the compound (B). Accordingly, the reaction is carried out at a desired temperature by changing these
10 conditions. Herein, the reaction temperature is preferably at most 200°C for enabling molding of the obtained composition, more preferably at most 180°C, and further preferably at most 150°C.

It is preferable to carry out processing at a temperature at which high molecular weight conversion or crosslinking does not occur
15 and processing can be carried out. But the temperature may be within a range in which processing can be carried out, though high molecular weight conversion occurs in part or partial crosslinking occurs.

In the case of the powder slash molding, it is preferable to
20 obtain the thermoplastic elastomer composition as powder. As the process of obtaining powder, the powder can be obtained by finely pulverizing the thermoplastic elastomer composition in a block state or a pellet state which is processed by the above-mentioned process by using an impact pulverizer such as a turbo mill, a pin mill, a hammer
25 mill and a centrifugation mill. At this time, pulverization is usually carried out at normal temperature, but mechanical pulverization can be carried out by using a cooling medium and a cooling facility.

<Molding process of thermoplastic elastomer composition>

The composition obtained in the above-described preparation process of the thermoplastic elastomer can be molded by various processes, and it can be preferably applied to powder slash molding, injection molding, injection blow molding, blow molding, extrusion blow molding, extrusion molding, calendar molding, vacuum molding, and press molding. Among them, the powder slash molding is more preferably used. Herein, the powder slash molding is a process of flowing the composition powder into a molding mold heated at a high temperature to be melt-molded and taking out a molded article solidified by cooling after taking a definite period of time. It is required to fluidize the powder to carry out melt-molding even under no pressurization in the powder slash molding, but on the other hand, the molded article after molding is exposed in environments where the molded article is used at 100°C or more. Therefore, it is difficult to make a balance between moldability and heat resistance. However, since the acrylic block copolymer (A) and the compound (B) in the thermoplastic elastomer composition of the present invention are in an unreacted state before molding, and are excellent in melting property in a mold. On the other hand, the acrylic block copolymer (A) is reacted with the compound (B) within a definite period of time until it is solidified by cooling, the acrylic block copolymer (A) is converted to having a high molecular weight or crosslinked, and heat resistance after the molding is improved. Consequently, the thermoplastic elastomer composition can be recognized as a preferable material for the powder slash molding.

When heat resistance is imparted by the high molecular

weight conversion of the acrylic block copolymer (A) at molding in the powder slash molding, the number average molecular weight of the acrylic block copolymer (A) after molding is preferably 100,000 or more, more preferably 150,000 or more and further preferably 200,000 or more. When the number average molecular weight is lower than 100,000, the improvement effect of the heat resistance tends to be lowered.

Further, when the heat resistance is imparted by crosslinking at molding, the insoluble content ratio (% by weight) of the molded article after molding is preferably 50 % or more by weight, more preferably 70 % or more by weight, and further more preferably 80 % or more by weight. When the insoluble content ratio is lower than 50 % by weight, the improvement effect of the moldability and heat resistance tends to be lowered. On the other hand, the insoluble content ratio before molding is preferably 30 % or less by weight, more preferably 10 % or less by weight, and further preferably 5 % or less by weight. When it is 30 % by weight or more, moldability tends to be deteriorated.

In the above description, the insoluble content ratio (% by weight) represents the weight of a residual solid for 1 g of the thermoplastic elastomer composition. The residual solid is obtained by packing 1 g of the thermoplastic elastomer composition in a 350 mesh metal net, immersing it in toluene at 80°C or acetone at 60°C for 24 hours (toluene or acetone is selected by the solubility of the thermoplastic elastomer composition), fractionating a toluene or acetone soluble part, drying the residual solid at 80°C under vacuum and measuring the weight g of the residual solid after drying.

EXAMPLES

The present invention is further explained in detail based on Examples, but the present invention is not limited only to these Examples.

5 Further, BA, EA, TBA, MMA, TBMA and GMA in Examples respectively represent n-butyl acrylate, ethyl acrylate, t-butyl acrylate, methyl methacrylate, t-butyl methacrylate and glycidyl methacrylate.
<Measurement method of molecular weight>

10 Molecular weight shown in the present Example was calculated in terms of polystyrene which was measured by a GPC analysis device shown below using chloroform as mobile phase. A GPC system made by Waters Co. was used as a system and Shodex k-804 (polystyrene gel) available from Showa Denko K.K. was used as a column.

15 <Measurement method of conversion of polymerization reaction>

The conversion of a polymerization reaction shown in the present Examples was measured by an analysis equipment and conditions shown below.

Device: Gas chromatography GC-14B made by Shimadzu Corporation.

20 Separation column: Capillary column; Supelcowax-10, 0.35 mm ϕ \times 30 m, made by J & W SCIENTIFIC INC.

Separation condition: Initial temperature: 60°C and retention for 3.5 minutes.

Temperature increasing speed: 40°C/min

25 Final temperature: 140°C and retention for 1.5 minutes

Injection temperature: 250°C

Detector temperature: 250°C

Sample adjustment: A sample was diluted to about 3-fold by ethyl acetate and butyl acetate was used as an internal standard substance.

<Measurement of insoluble content ratio (% by weight)>

1 g of the thermoplastic elastomer composition was packed
5 in a 350 mesh metal net, and it was immersed in toluene at 80°C or acetone at 60°C for 24 hours (toluene or acetone is selected by the solubility of the acrylic block copolymer). Then, a toluene or acetone soluble part was fractionated, the residual solid was dried at 80°C under vacuum and the weight g of the residual solid after drying was
10 measured. The insoluble content ratio is represented by a weight of the residual solid content for 1 g of the thermoplastic elastomer composition.

<Ethanol resistance test>

Ethanol resistance shown in Examples and Comparative
15 Examples was measured according to conditions shown in the following.

The sheets with crepe pattern prepared in Examples and Comparative Examples were placed on a flat surface, one drop of ethanol (available from Wako Pure Chemical Industry Ltd.) was added
20 dropwise with a pipette thereon, and they were left alone at a room temperature for 24 hours. Then, their surfaces were visually observed. The sheets were evaluated by the criteria such that those having no trace were ○, those in which trace was observed but had no whitening were △, and those in which whitening was observed were ×.

25 <Oil resistance test>

Paraffin resistance property shown in Examples and Comparative Examples were measured by conditions shown in the

following.

The sheets with crepe pattern prepared in Examples and Comparative Examples were placed on a flat surface, one drop of liquid paraffin (available from Nakarai Tesk Co.) was added dropwise with a
5 pipette thereon, and they were left alone at a room temperature for 24 hours. Then, the liquid paraffin was wiped off with KIMWIPE (made by Cresia Inc.) and their surfaces were visually observed. The sheets were evaluated by the criteria such that those having no trace were ○, those in which trace was observed but had no whitening were △, and
10 those in which whitening was observed were ×.

<Heat resistance test>

Heat resistance shown in Examples and Comparative Examples were measured by conditions shown in the following.

The sheets with crepe pattern prepared in Examples and
15 Comparative Examples were left alone at 120°C for 24 hours. Then, their surfaces were visually observed. The sheets were evaluated by the criteria such that those in which the change of the crepe pattern was not observed were ○, those in which the change of the crepe pattern was not clear but surface luster was increased in comparison
20 with the initial stage were △, and those in which the change of the crepe pattern was observed were ×.

<Urethane adhesiveness test>

Superficial skin materials were prepared by press-molding the compositions according to Examples and Comparative Examples.
25 A cartridge type polyurethane (available from Air Tight Co.) in which 4,4'-diphenylmethanediisocyanate was the main component was coated on a metal plate and the superficial skin material was

immediately mounted on the foamed body to be adhered. After taking for at least 12 hours (state in which the foamed article was completely cured), the superficial skin material was peeled from the foamed urethane with a hand and the state of destruction was observed.

- 5 Those in which destruction occurred in a urethane material were referred to as ○, those in which destruction occurred at the portion of interface between a sheet and urethane were referred to as △, and those in which destruction occurred at interface between the sheet and urethane were referred to as ×.

10 <Powder slash property test>

A block of the composition was prepared according to Examples and Comparative Examples. The block of the composition was charged in a small size pulverizer (made by Kyoritsu-Riko Co.) cooled with dry ice and pulverized while adding dry ice. The obtained
15 powder was evaluated under condition in the following. The obtained powder was brought in contact for 30 seconds with a skin crepe metal plate heated at 260°C, unmelted powder was removed after heat-melting and it was cooled to a room temperature to obtain a molded sheet.

20 Evaluation index:

The obtained molded sheet was observed visually. The sheet was evaluated by the criteria such that those in which crepe transcription property was good and pinhole/foams were not found were referred to as ○, those in which either of one of items was poor
25 were referred to as △, and those in which the site of poor crepe formation was found and pinhole/foams were found were referred to as ×.

PREPARATION EXAMPLE 1

Synthesis of (MMA-co-BA-co-TBMA)-b-BA-b-(MMA-co-BA-co-TBMA) type acrylic block copolymer (hereinafter, described as "the precursor 1")

5 Following operations were carried out in order to obtain the precursor 1. After the atmosphere of a 15 L pressure resistant reactor was purged with nitrogen, 13.6 g (95 mmol) of copper bromide was weighed and 146 g of acetonitrile (treated with nitrogen bubbling) was added. After stirring by heating at 70°C for 30 minutes, 19.0 g
10 (53 mmol) of diethyl 2,5-dibromoadipate as an initiator and 1664 g (13.0 mol) of BA were added. The mixture was stirred by heating at 85°C and 1.65 g (9.5 mmol) of pentamethyldiethylenetriamine as a ligand was added to initiate polymerization.

 About 0.2 mL of the polymerization solution as for
15 sampling was extracted from the polymerization solution periodically from the start of polymerization and the conversion of BA was determined by the gas chromatogram analysis of the sampling solution. Polymerization speed was controlled by adding pentamethyldiethylenetriamine as needed. When the conversion of BA
20 reached at 94.6 %, 82.8 g (0.58 mol) of TBMA, 927 g (9.3 mol) of MMA, 202 g (1.6 mol) of BA, 9.4 g (95 mmol) of copper chloride, 1.98 g (9.5 mmol) of pentamethyldiethylenetriamine and 2269 g of toluene (treated with nitrogen bubbling) were added thereto. Similarly, the conversion of TBMA, MMA and BA were determined. When the conversion of
25 TBMA was 93.1 %, the conversion of MMA was 89.2 % and the conversion of BA was 66.8 % based on the concentration of BA just after the addition of MMA/TBMA/BA, 2400 g of toluene was added,

and the reactor was cooled with a water bath to terminate the reaction.

The polymerization solution was diluted with 1900 g of toluene, 32.5 g of p-toluene sulfonic acid monohydrate was added to be stirred for 3 hours at a room temperature, and a solid precipitated was removed by filtration. 40.8 g of an absorbent KYOWARD 500SH (available from KYOWA Chemical Industry Co., Ltd.) was added to the obtained polymer solution and the mixture was further stirred at a room temperature for one hour. The absorbent was filtered by a Kiriyaama funnel to obtain a colorless transparent polymer solution. The solution was dried and the solvent and residual monomer were removed to obtain the precursor 1 which was the objective acrylic block copolymer.

When the GPC analysis of the precursor 1 of the acrylic block copolymer was carried out, the number average molecular weight M_n was 72,200 and the molecular weight distribution M_w/M_n was 1.42.

<Reaction forming acid anhydride of precursor 1>

45 g of the precursor 1 obtained in the above description and 0.09 g of Irganox 1010 (available from Chiba Specialty Chemicals Co., Ltd.) were melt-kneaded at 100 rpm for 20 minutes using a LABO Plasto Mill 50C150 (blade shape: roller type R60, made by TOYO SEIKI KOGYO Co., Ltd.) set at 240°C to obtain an objective acrylic block copolymer containing an acid anhydride group and a carboxyl group (hereinafter, the obtained polymer is described as "the polymer 1").

The conversion to the acid anhydride group and the carboxyl group at a t-butyl ester site could be confirmed by IR (infrared absorption spectrum) and $^{13}\text{C}(^1\text{H})$ -NMR (nuclear magnetic resonance

spectrum). Namely, in IR, it could be confirmed because absorption spectrum derived from an acid anhydride group was observed nearby 1800 cm^{-1} after the conversion. In $^{13}\text{C}(^1\text{H})$ -NMR, it could be confirmed because a signal at 82 ppm which was derived from quaternary carbon of t-butyl group and a signal at 28 ppm which was derived from methyl carbon of t-butyl group were extinguished after the conversion and a signal at 172 to 173 ppm (m) which was derived from the carbonyl carbon of an acid anhydride group and a signal at 176 to 179 ppm (m) which was derived from the carbonyl carbon of a carboxyl group newly appeared.

PREPARATION EXAMPLE 2

Synthesis of (MMA-co-EA)-b-(BA-co-TBA)-b-(MMA-co-EA) type acrylic block copolymer (hereinafter, described as "the precursor 2")

The following operations were carried out in order to obtain the precursor 2.

A 500 L reactor purged with nitrogen was charged with 71.83 kg of n-butyl acrylate, 3.23 kg of t-butyl acrylate and 0.804 kg of cuprous bromide to initiate stirring. Then, a solution in which 1.12 kg of diethyl 2,5-dibromoadipate was dissolved in 6.59 kg of acetonitrile was charged thereto and the inner solution was stirred for 30 minutes while flowing warm water into jacket and raising the temperature of inner solution to 75°C. When the inner temperature reached at 75°C, 97.1 g of pentamethyldiethylenetriamine was added to initiate the polymerization of an acrylic polymer block. Polymerization speed was controlled by adding pentamethyldiethylenetriamine at polymerization as needed.

Pentamethyldiethylenetriamine was added 3 times in total (291.3 g in total) during the polymerization of the acrylic polymer block.

When the conversion reached at 99.1 %, 97.6 kg of toluene, 0.555 kg of cuprous chloride, 45.29 kg of methyl methacrylate, 7.36 kg of ethyl acrylate and 97.1 g of pentamethyldiethylenetriamine were added thereto to start the polymerization of the methacrylic polymer block. When the conversion of methyl methacrylate reached at 95.8 %, 220 kg of toluene was added to dilute the reaction solution and the reactor was cooled to terminate the reaction. When the GPC analysis of the obtained block copolymer was carried out, the number average molecular weight M_n was 62400 and the molecular weight distribution M_w/M_n was 1.44.

30 kg of toluene was added to the obtained block copolymer solution, and a concentration of the polymer was adjusted to 25 % by weight. To the solution, 2.24 kg of p-toluene sulfonic acid was added, the reactor was purged with nitrogen, and the solution was stirred for 3 hours at 30°C. The reaction solution was sampled and after confirming that the solution was colorless and transparent, 2.44 kg of RADIOLITE #3000 (available from Showa Chemical Industry Co., Ltd.) was added to the solution. Then, the reactor was pressurized to 0.1 to 0.4 MPaG with nitrogen and solid was separated using a pressurized filter (filtration area of 0.45 m²) equipped with polyester felt as a filtration material.

2.44 kg of KYOWARD 500SH was added to about 450 kg of the block copolymer solution after filtration, the reactor was purged with nitrogen and the solution was stirred at 30°C for one hour. The reaction solution was sampled and after confirming that the solution

was neutral, the reaction was terminated. Then, the reactor was pressurized to 0.1 to 0.4 MPaG with nitrogen and solid was separated using a pressurized filter (filtration area of 0.45 m²) equipped with polyester felt as a filtration material to obtain the polymer solution.

5 728 g of Irganox 1010 (available from Chiba Specialty Chemicals Co., Ltd.) was added to the obtained polymer solution and dissolved.

Successively, the solvent component was evaporated from the polymer solution. SCP 100 (heat transfer area: 1 m²) made by Kurimoto Ltd. was used as an evaporator. The evaporation of the polymer solution was carried out by setting heat transfer oil at the inlet

10 of the evaporator at 180°C, the vacuum degree of the evaporator as 90 Torr, the rotational number of a screw as 60 rpm and the feed speed of the polymer solution as 32 kg/h. The polymer was made as strand with a dice of 4 mmφ through a discharger, it was cooled with a water

15 vessel filled with 3 % suspension solution of ALFLOW H50ES (main component: ethylenebisstearic acid amide available from NOF Corporation) and then, columnar pellets were obtained by a pelletizer. Thus, pellets of the precursor 2 of the objective acrylic block copolymer were obtained.

20 <Reaction forming acid anhydride of precursor 2>

1 part by weight of hydrotalcite DHT-4A-2 (available from KYOWA Chemical Industry Co., Ltd.) as an acid trapping agent was compounded to 100 parts by weight of the precursor 2 obtained in the above description, and extrusion-kneaded by setting a rotational

25 number at 150 rpm, the cylinder temperature of a hopper setting portion at 100°C and all other setting temperature at 260°C using a biaxial extruder with a vent (44 mm, L/D = 42.25) (made by Japan

Steel Works Ltd.) to obtain an objective acrylic block copolymer containing an acid anhydride group and a carboxyl group (hereinafter, the obtained polymer is described as "the polymer 2"). At extrusion, the vent orifice was blocked. Further, at this time, an underwater cut
5 pelletizer (CLS-6-8.1 COMPACT LAB SYSTEM, made by GALA INDUSTRIES INC.) was connected with the edge of the biaxial extruder and spherical pellets without adhesion preventing property were obtained by adding ALFLOW H-50ES (available from NOF Corporation) as an adhesion preventing agent in the circulation water of the
10 underwater cut pelletizer.

The measurement of conversion efficiency to the acid anhydride group and the carboxyl group at a t-butyl ester site was carried out by quantifying the amount of isobutylene generated from the t-butyl group by a thermal decomposition reaction at 280°C. As a
15 result of the measurement, the conversion efficiency of the obtained resin was at least 95 %.

PREPARATION EXAMPLE 3

Synthesis of (MMA-co-EA)-b-(BA-co-GMA)-b-(MMA-co-EA) type acrylic
20 block copolymer (hereinafter, described as "the polymer 3")

The following operations were carried out in order to obtain the precursor 3. After a 15 L pressure resistant reactor was purged with nitrogen, 6.45 g (45 mmol) of copper bromide, 712 g (5.6 mol) of BA and 54 g (0.38 mol) of GMA were charged thereto to initiate stirring.
25 Then, a solution in which 9.0 g (25 mmol) of diethyl 2,5-dibromoadipate as an initiator was dissolved in 67 g of acetonitrile (treated with nitrogen bubbling) was charged thereto and the inner

solution was stirred for 30 minutes while raising the temperature of the inner solution to 75°C. When the inner temperature reached at 75°C, 0.78 g (4 mmol) of pentamethyldiethylenetriamine as a ligand was added to initiate polymerization of an acrylic polymer block.

5 About 0.2 mL of the polymerization solution as for sampling was extracted from the polymerization solution periodically from the start of the polymerization and the conversions of BA and GMA were determined by the gas chromatogram analysis of the sampling solution. Polymerization speed was controlled by adding
10 pentamethyldiethylenetriamine at polymerization as needed. When the conversion of BA reached at 96.0 % and the conversion of GMA reached at 100 %, 447 g (4.5 mol) of MMA, 124 g (1.24 mol) of EA, 4.45 g (45 mmol) of copper chloride, 0.78 g (4 mmol) of pentamethyldiethylenetriamine and 1059 g of toluene (treated with
15 nitrogen bubbling) were added thereto to initiate polymerization of the methacrylic polymer block.

 The conversion of MMA were determined in the same manner as at the polymerization of the acrylic polymer block. Sampling was carried out at charging MMA and the conversion of MMA
20 was determined using this as a basis. After charging MMA, the inner temperature was set at 85°C. A polymerization speed was controlled by adding pentamethyldiethylenetriamine at polymerization as needed. When the conversion of MMA was 90.0 %, 1060 g of toluene was added and the reactor was cooled with a water bath to terminate the reaction.

25 Toluene was added to the obtained reaction solution and it was diluted so that a concentration of the polymer was 25 % by weight. 15.4 g of p-toluene sulfonic acid monohydrate was added to the

solution to be stirred for 3 hours at a room temperature, and a precipitated solid content was removed by filtration.

18.5 g of an absorbent KYOWARD 500SH (available from KYOWA Chemical Industry Co., Ltd.) was added to the obtained polymer solution, and the mixture was further stirred at a room temperature for one hour. The absorbent was filtered by a Kiriya
5 funnel to obtain a colorless transparent polymer solution. The solution was dried and the solvent and the residual monomer were removed to obtain the polymer 3 which was the objective acrylic block
10 copolymer.

When the GPC analysis of the polymer 3 of the acrylic block copolymer was carried out, the number average molecular weight M_n was 72,992, and the molecular weight distribution M_w/M_n was 1.40.

15

PREPARATION EXAMPLE 4

Synthesis of (MMA-co-BA)-b-BA-b-(MMA-co-BA) type acrylic block copolymer (hereinafter, described as the polymer 4)

The following operations were carried out for obtaining the polymer 4. After a 5 L separable flask as a polymerization container was purged with nitrogen, 5.7 g (40 mmol) of copper bromide was weighed and 59 g of acetonitrile (treated with nitrogen bubbling) was added thereto. After stirring by heating at 70°C for 30 minutes, 8.0 g (22 mmol) of diethyl 2,5-dibromoadipate as an initiator and 671 g (5.2
20 mol) of BA were added thereto. The mixture was stirred by heating at 85°C and 0.69 g (4.0 mmol) of pentamethyldiethylenetriamine as a
25 ligand was added to initiate polymerization.

About 0.2 mL of polymerization solution as for sampling was extracted from the polymerization solution periodically from the initiation of polymerization and the conversion of BA was determined by the gas chromatogram analysis of the sampling solution.

5 Polymerization speed was controlled by adding pentamethyldiethylenetriamine as needed. When the conversion of BA reached at 94.5 %, 391 g (3.9 mol) of MMA, 61 g (0.47 mol) of BA, 4.0 g (40 mmol) of copper chloride, 0.69 g (4.0 mmol) of pentamethyldiethylenetriamine and 841 g of toluene (treated with

10 nitrogen bubbling) were added thereto. Similarly, the conversions of MMA and BA were determined. When the conversion of MMA was 90.5 % and the conversion of BA was 67.1 % based on the concentration of BA just after the addition of MMA/TBMA/BA, 1500 g of toluene was added thereto and the reactor was cooled with a water

15 bath to terminate the reaction.

The reaction solution was diluted with 700 g of toluene, 16.0 g of p-toluene sulfonic acid monohydrate was added to be stirred for 3 hours at a room temperature, and a precipitated solid content was removed by filtration. 16.0 g of an absorbent KYOWARD 500SH

20 (available from KYOWA Chemical Industry Co., Ltd.) was added to the obtained polymer solution, and the mixture was further stirred at a room temperature for one hour. The absorbent was filtered by a Kiriyaama funnel to obtain a colorless transparent polymer solution. The solution was dried, and the solvent and the residual monomer

25 were removed to obtain the objective polymer 4.

When the GPC analysis of the obtained polymer 4 was carried out, the number average molecular weight M_n was 62600 and

the molecular weight distribution M_w/M_n was 1.44.

PREPARATION EXAMPLE 5

Synthesis of (MMA-co-BA)-b-BA-b-(MMA-co-BA) type acrylic block
5 copolymer (hereinafter, described as the polymer 5)

The following operations were carried out in order to obtain the polymer 5.

After a 5 l separable flask as a polymerization container was purged with nitrogen, 11.3 g (78.5 mmol) of copper bromide was
10 weighed and 180 ml of acetonitrile (dried with molecular sieves 3A and then treated with nitrogen bubbling) was added thereto. After stirring by heating at 70°C for 5 minutes, it was cooled to a room temperature again, 5.7 g (15.7 mmol) of diethyl 2,5-dibromoadipate as an initiator and 804.6 g (900.0 ml) of n-butyl acrylate were added. The mixture
15 was stirred by heating at 80°C and 1.6 ml (7.9 mmol) of pentamethyldiethylenetriamine as a ligand was added to initiate polymerization. About 0.2 ml of a polymerization solution as for sampling was extracted from the polymerization solution periodically from the initiation of polymerization and the conversion of butyl
20 acrylate was determined by the gas chromatogram analysis of the sampling solution. Polymerization speed was controlled by adding pentamethyldiethylenetriamine as needed. When the conversion of n-butyl acrylate reached at 95 %, 345.7 g (369.3 ml) of methyl methacrylate, 7.8 g (78.5 mmol) of copper chloride, 1.6 ml (7.9 mmol)
25 of pentamethyldiethylenetriamine and 1107.9 ml of toluene (dried with molecular sieves 3A and then treated with nitrogen bubbling) were added thereto. Similarly, the conversion of methyl methacrylate was

determined. When the conversion of methyl methacrylate was 85 % and the conversion of n-butyl acrylate was 98 %, 1500 ml of toluene was added and the reactor was cooled with a water bath to terminate the reaction. The polymerization solution was always green during the
5 reaction.

The reaction solution was diluted with 4000 ml of toluene, 22.1 g of p-toluene sulfonic acid monohydrate was added to be stirred at 23°C for 3 hours. After removing the insoluble portion precipitated by filtering with a Kiriama funnel, 9.7 g of an absorbent KYOWARD
10 500SH was added to the polymer solution and the mixture was further stirred at 23°C for 3 hours. The absorbent was filtered by a Kiriama funnel to obtain a colorless transparent polymer solution. The solution was dried, and the solvent and the residual monomer were removed to obtain the objective polymer 5.

15 When the GPC analysis of the obtained polymer 5 was carried out, the number average molecular weight M_n was 119,200 and the molecular weight distribution M_w/M_n was 1.51. Further, when compositional analysis by NMR was carried out, it was BA/MMA = 72/28 (% by weight).

20

EXAMPLE 1

A proportion of 10 parts by weight of EPICOAT 828 (available from Japan Epoxy Resins Co., Ltd.) which is an epoxy resin, 0.5 part by weight of carbon black (ASAHI #15; available from Asahi
25 Carbon Co., Ltd.) and 0.3 part by weight of Irganox 1010 (available from Chiba Specialty Chemicals Co., Ltd.) based on 100 parts by weight (38 g) of the polymer 1 obtained in Preparation example 1 was

melt-kneaded at 100 rpm for 15 minutes using a LABO Plasto Mill 50C150 (blade shape: roller type R60, made by TOYO SEIKI KOGYO Co., Ltd.) set at 100°C to obtain a block sample.

The obtained sample was molded at a set temperature of 200°C for 5 minutes with a hot press (a compression molding machine NSF-50 made by Shinto Metal Industries Ltd.) using a skin crepe metal plate and molded articles for evaluation having a thickness of 1mm on which the skin crepe pattern was transcribed were obtained. Ethanol resistance, oil resistance, urethane adhesivity and heat resistance test were measured for these molded articles. The result is shown in Table 1. Further, the powder slash property test was carried out for powder obtained by pulverizing the block sample obtained in the above description. Further, an insoluble content ratio (% by weight) was measured by using a powder before the powder slash molding and a sheet after molding. The result is shown in Table 1.

EXAMPLE 2

A proportion of 4.4 parts by weight of EPIOL E-400 (available from NOF Corporation) which is an epoxy group-containing compound and 0.5 part by weight of carbon black (ASAHI #15; available from Asahi Carbon Co., Ltd.) based on 100 parts by weight (38 g) of the polymer 2 obtained in Preparation Example 2 was melt-kneaded at 100 rpm for 15 minutes using a LABO Plasto Mill 50C150 (blade shape: roller type R60, made by TOYO SEIKI KOGYO Co., Ltd.) set at 100°C to obtain a block sample.

The obtained sample was evaluated in the same manner as Example 1. The result is shown in Table 1.

EXAMPLE 3

The operation was carried out in the same manner as Example 2 except that 2.5 parts by weight of EPIOL P-200 (available from NOF Corporation) was used in place of 4.4 parts by weight of EPIOL E-400 (available from NOF Corporation) of Example 2, and a block sample was obtained to be evaluated.

EXAMPLE 4.

A proportion of 2.0 parts by weight of triethylene glycol (available from Wako Pure Chemical Industry Ltd.) which is a hydroxyl group-containing compound, 0.5 part by weight of carbon black (ASAHI #15; available from Asahi Carbon Co., Ltd.) and 0.3 part by weight of Irganox 1010 (available from Chiba Specialty Chemicals Co., Ltd.) based on 100 parts by weight (38 g) of the polymer 3 obtained in Preparation Example 3 was melt-kneaded at 100 rpm for 15 minutes using a LABO Plasto Mill 50C150 (blade shape: roller type R60, made by TOYO SEIKI KOGYO Co., Ltd.) set at 100°C to obtain a block sample.

The obtained sample was evaluated in the same manner as Example 1. The result is shown in Table 1.

COMPARATIVE EXAMPLE 1

A proportion of 0.5 part by weight of carbon black (ASAHI #15; available from Asahi Carbon Co., Ltd.) and 0.3 part by weight of Irganox 1010 (available from Chiba Specialty Chemicals Co. Ltd.) based on 100 parts by weight (38 g) of the polymer 4 obtained in Preparation example 4 was melt-kneaded at 100 rpm for 15 minutes using a LABO

Plasto Mill 50C150 (blade shape: roller type R60, made by TOYO SEIKI KOGYO Co., Ltd.) set at 190°C to obtain a block sample.

The obtained sample was evaluated in the same manner as Example 1. The result is shown in Table 1. It is clear that the block sample is excellent in powder slash moldability but inferior in heat resistance.

COMPARATIVE EXAMPLE 2

The operation was carried out in the same manner as Comparative Example 1 except that the polymer 5 was used in place of the polymer 4 of Comparative Example 1, and a block sample was obtained to be evaluated.

It is clear that the block sample is excellent in heat resistance but inferior in powder slash moldability.

TABLE 1

	Ex.				Com. Ex.	
	1	2	3	4	1	2
Polymer 1	100					
Polymer 2		100	100			
Polymer 3				100		
Polymer 4					100	
Polymer 5						100
EPICOAT 828	10					
EPIOL E-400		4.4				
EPIOL P-200			2.5			
Triethylene glycol				2		
ASAHI #15	0.5	0.5	0.5	0.5	0.5	0.5
Ethanol resistance	○	○	○	○	○	○
Oil resistance	○	○	○	○	○	○
Heat resistance	○	○	○	○	×	○
Urethane adhesivity	○	○	○	○	△	△
Powder slash property	○	○	○	○	○	×
Insoluble content ratio before molding (% by weight)	0	-	-	-	0	0
Insoluble content ratio after molding (% by weight)	64	-	-	-	0	0

As cleared from Table 1 (Examples 1 to 4 and Comparative Examples 1 and 2), it proves that the thermoplastic elastomer composition of the present invention is excellent in powder slash moldability and, additionally, improves heat resistance of the obtained molded article by a crosslinking reaction (the increase of insoluble content ratio) compared with only a block copolymer. Further, it is clear that it is also excellent in ethanol resistance and oil resistance.

Further, when the obtained sheet is used as a superficial skin material for an automobile, it is required to be adhered to polyurethane which are generally used as a substrate, and it is clear that the thermoplastic

elastomer composition is favorably adhered.

INDUSTRIAL APPLICABILITY

Since the molded article obtained by molding the
5 thermoplastic elastomer composition of the present invention is
excellent in heat resistance, weather resistance, chemical resistance,
adhesivity, flexibility, abrasion resistance, it can be used as materials
having purposes such as a superficial material, a touch material, an
appearance material, an abrasive resistant material, an oil resistant
10 material, a dumping material and an adherent material; as arbitrary
shapes such as a sheet, a flat panel, a film, a small size molded article,
a large size molded article and the other shapes; further, as parts such
as panels, steerings, grips and switches; and further as a sealing
material. Their uses are not particularly limited, but uses for an
15 automobile, for home electronics products or for electric products for
office supplies are exemplified. Examples thereof are a superficial
skin material for an automobile, a touch material for an automobile, an
appearance material for an automobile, panels for an automobile,
steerings for an automobile, grips for an automobile, switches for an
20 automobile and panels for electric products at home or office, switches
for electric products at home or office. Among those, it is preferably
used for a superficial skin for an automobile interior.